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(71) Applicant (for all designated States except US): INCYTE GENOMICS, INC. [US/US]; 3160 Porter Drive, Palo Alto, CA 94304 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): HODGSON, David, M. [US/US]; 567 Addison Avenue, Palo Alto, CA 94304 (US). LINCOLN, Stephen, E. [US/US]; 725 Sapphire Street, Redwood City, CA 94061 (US). RUSSO, Frank, D. [US/US]; 1583 Courdillaeras Road, Redwood City, CA 94062 (US). SPIRO, Peter, A. [US/US]; 3875 Park Boulevard, Apt. B16, Palo Alto, CA 94306 (US). BANVILLE,

Steven, C. [US/US]; 604 San Diego Avenue, Sunnyvale, CA 94086 (US). BRATCHER, Shawn, R. [US/US]; 550 Ortega Avenue #B321, Mountain View, CA 94040 (US). DUFOUR, Gerard, E. [US/US]; 5327 Greenridge Road, Castro Valley, CA 94552-2619 (US). COHEN, Howard, J. [US/US]; 3272 Cowper Street, Palo Alto, CA 94306-3004 (US). ROSEN, Bruce, H. [US/US]; 177 Hanna Way, Menlo Park, CA 94025 (US). SHAH, Purvi [IN/US]; 859 Salt Lake Drive, San Jose, CA 95133 (US). CHALUP, Michael, S. [US/US]; 183 Acalanes Drive, Apt. 6, Sunnyvale, CA 94086 (US). HILLMAN, Jennifer, L. [US/US]; 230 Monroe Drive, #17, Mountain View, CA 94040 (US). JONES, Anissa, Lee [US/US]; 445 South 15th Street, San Jose, CA 95112 (US). YU, Jimmy, Y. [US/US]; 37330 Portico Terrace, Fremont, CA 94536-7901 (US). GREENAWALT, Lila, B. [US/US]; 1596 Ballantree Way, San Jose, CA 95118-2106 (US). PANZER, Scott, R. [US/US]; 965 East El Camino, #621, Sunnyvale, CA 94087 (US). ROSEBERRY, Ann, M. [US/US]; 725 Sapphire Street, Redwood City, CA 94061 (US). WRIGHT, Rachel, J. [NZ/US]; 339 Anna Way, Mountain View, CA 94043 (US). CHEN, Wensheng [CN/US]; 210 Easy Street, #25, Mountain View, CA 94043 (US). LIU, Tommy, F. [US/US]; 201 Ottilia Street, Daly City, Ca 94014 (US). YAP, Pierre, E. [US/US]; 201 Happy Hollow Court, Lafayette, CA 94549-6243 (US). STOCKDREHER, Theresa, K. [US/US]; 1596 Ontario Drive, #2, Sunnyvale, CA 94087 (US). AMSHEY, Stefan [US/US]; 1541 Canna Court, Mountain View, CA 94043 (US). FONG, Willy, T. [US/US]; 572 Cambridge Street, San Francisco, CA 94134 (US).

- (74) Agents: HAMLET-COX, Diana et al.; Incyte Genomics, Inc., 3160 Porter Drive, Palo Alto, CA 94304 (US).
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(54) Title: SECRETORY MOLECULES

(57) Abstract: The present invention provides purified secretory polynucleotides (sptm). Also encompassed are the polypeptides (SPTM) encoded by sptm. The invention also provides for the use of sptm, or complements, oligonucleotides, or fragments thereof in diagnostic assays. The invention further provides for vectors and host cells containing sptm for the expression of SPTM. The invention additionally provides for the use of isolated and purified SPTM to induce antibodies and to screen libraries of compounds and the use of anti-SPTM antibodies in diagnostic assays. Also provided are microarrays containing sptm and methods of use.



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SECRETORY MOLECULES

TECHNICAL FIELD

The present invention relates to secretory molecules and to the use of these sequences in the diagnosis, study, prevention, and treatment of diseases associated with, as well as effects of exogenous compounds on, cell signaling and the expression of secretory molecules.

BACKGROUND OF THE INVENTION

Protein transport and secretion are essential for cellular function. Protein transport is mediated by a signal peptide located at the amino terminus of the protein to be transported or secreted. The signal peptide is comprised of about ten to twenty hydrophobic amino acids which target the nascent protein from the ribosome to a particular membrane bound compartment such as the endoplasmic reticulum (ER). Proteins targeted to the ER may either proceed through the secretory pathway or remain in any of the secretory organelles such as the ER, Golgi apparatus, or lysosomes. Proteins that transit through the secretory pathway are either secreted into the extracellular space or retained in the plasma membrane. Proteins that are retained in the plasma membrane contain one or more transmembrane domains, each comprised of about 20 hydrophobic amino acid residues. Proteins that are secreted from the cell are generally synthesized as inactive precursors that are activated by posttranslational processing events during transit through the secretory pathway. Such events include glycosylation, proteolysis, and removal of the signal peptide by a signal peptidase. Other events that may occur during protein transport include chaperone-dependent unfolding and folding of the nascent protein and interaction of the protein with a receptor or pore complex. Examples of secretory proteins with amino terminal signal peptides are discussed below and include proteins with important roles in cell-to-cell signaling. Such proteins include transmembrane receptors and cell surface markers, extracellular matrix molecules, cytokines, hormones, growth and differentiation factors, neuropeptides, vasomediators, ion channels, transporters/pumps, and proteases. (Reviewed in Alberts, B. et al. (1994) Molecular Biology of The Cell, Garland Publishing, New York, NY, pp. 557-560, 582-592.)

G-protein coupled receptors (GPCRs) comprise a superfamily of integral membrane proteins which transduce extracellular signals. Not all GPCRs contain N-terminal signal peptides. GPCRs include receptors for biogenic amines such as dopamine, epinephrine, histamine, glutamate (metabotropic-type), acetylcholine (muscarinic-type), and scrotonin; for lipid mediators of inflammation such as prostaglandins, platelet activating factor, and leukotrienes; for peptide hormones such as calcitonin, C5a anaphylatoxin, follicle stimulating hormone, gonadotropin releasing hormone, neurokinin, oxytocin, and thrombin; and for sensory signal mediators such as retinal photopigments and olfactory stimulatory molecules. The structure of these highly conserved receptors consists of seven

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hydrophobic transmembrane regions, cysteine disulfide bridges between the second and third extracellular loops, an extracellular N-terminus, and a cytoplasmic C-terminus. The N-terminus interacts with ligands, the disulfide bridges interact with agonists and antagonists, and the large third intracellular loop interacts with G proteins to activate second messengers such as cyclic AMP, phospholipase C, inositol triphosphate, or ion channels. (Reviewed in Watson, S. and Arkinstall, S. (1994) The G-protein Linked Receptor Facts Book, Academic Press, San Diego, CA, pp. 2-6; and Bolander, F.F. (1994) Molecular Endocrinology, Academic Press, San Diego, CA, pp. 162-176.)

Other types of receptors include cell surface antigens identified on leukocytic cells of the immune system. These antigens have been identified using systematic, monoclonal antibody (mAb)based "shot gun" techniques. These techniques have resulted in the production of hundreds of mAbs 10 directed against unknown cell surface leukocytic antigens. These antigens have been grouped into "clusters of differentiation" based on common immunocytochemical localization patterns in various differentiated and undifferentiated leukocytic cell types. Antigens in a given cluster are presumed to identify a single cell surface protein and are assigned a "cluster of differentiation" or "CD" designation. Some of the genes encoding proteins identified by CD antigens have been cloned and 15 verified by standard molecular biology techniques. CD antigens have been characterized as both transmembrane proteins and cell surface proteins anchored to the plasma membrane via covalent attachment to fatty acid-containing glycolipids such as glycosylphosphatidylinositol (GPI). (Reviewed in Barclay, A. N. et al. (1995) The Leucocyte Antigen Facts Book, Academic Press, San Diego, CA, pp. 17-20.)

Matrix proteins (MPs) are transmembrane and extracellular proteins which function in formation, growth, remodeling, and maintenance of tissues and as important mediators and regulators of the inflammatory response. The expression and balance of MPs may be perturbed by biochemical changes that result from congenital, epigenetic, or infectious diseases. In addition, MPs affect leukocyte migration, proliferation, differentiation, and activation in the immune response. MPs are 25 frequently characterized by the presence of one or more domains which may include collagen-like domains, EGF-like domains, immunoglobulin-like domains, and fibronectin-like domains. In addition, MPs may be heavily glycosylated and may contain an Arginine-Glycine-Aspartate (RGD) tripeptide motif which may play a role in adhesive interactions. MPs include extracellular proteins such as fibronectin, collagen, galectin, vitronectin and its proteolytic derivative somatomedin B; and cell adhesion receptors such as cell adhesion molecules (CAMs), cadherins, and integrins. (Reviewed in Ayad, S. et al. (1994) The Extracellular Matrix Facts Book, Academic Press, San Diego, CA, pp. 2-16; Ruoslahti, E. (1997) Kidney Int. 51:1413-1417; Sjaastad, M.D. and Nelson, W.J. (1997) BioEssays 35

Cytokines are secreted by hematopoietic cells in response to injury or infection. Interleukins,

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Ion channels, ion pumps, and transport proteins mediate the transport of molecules across cellular membranes. Transport can occur by a passive, concentration-dependent mechanism or can be linked to an energy source such as ATP hydrolysis. Symporters and antiporters transport ions and small molecules such as amino acids, glucose, and drugs. Symporters transport molecules and ions unidirectionally, and antiporters transport molecules and ions bidirectionally. Transporter superfamilies include facilitative transporters and active ATP-binding cassette transporters which are involved in multiple-drug resistance and the targeting of antigenic peptides to MHC Class I molecules. These transporters bind to a specific ion or other molecule and undergo a conformational change in order to transfer the ion or molecule across the membrane. (Reviewed in Alberts, B. et al. (1994) Molecular Biology of The Cell, Garland Publishing, New York, NY, pp. 523-546.)

Ion channels are formed by transmembrane proteins which create a lined passageway across the membrane through which water and ions, such as Na⁺, K⁺, Ca²⁺, and Cl⁻, enter and exit the cell. For example, chloride channels are involved in the regulation of the membrane electric potential as well as absorption and secretion of ions across the membrane. Chloride channels also regulate the internal pH of membrane-bound organelles.

Ion pumps are ATPases which actively maintain membrane gradients. Ion pumps are classified as P, V, or F according to their structure and function. All have one or more binding sites for ATP in their cytosolic domains. The P-class ion pumps include Ca^{2+} ATPase and Na^+/K^+ ATPase and function in transporting H^+ , Na^+ , K^+ , and Ca^{2+} ions. P-class pumps consist of two α and two β transmembrane subunits. The V- and F-class ion pumps have similar structures but transport only H^+ . F class H^+ pumps mediate transport across the membranes of mitochondria and chloroplasts, while V-class H^+ pumps regulate acidity inside lysosomes, endosomes, and plant vacuoles.

A family of structurally related intrinsic membrane proteins known as facilitative glucose transporters catalyze the movement of glucose and other selected sugars across the plasma membrane. The proteins in this family contain a highly conserved, large transmembrane domain comprised of 12 α -helices, and several weakly conserved, cytoplasmic and exoplasmic domains. (Pessin, J. E., and Bell, G.I. (1992) Annu. Rev. Physiol. 54:911-930.)

Amino acid transport is mediated by Na⁺ dependent amino acid transporters. These transporters are involved in gastrointestinal and renal uptake of dietary and cellular amino acids and in neuronal reuptake of neurotransmitters. Transport of cationic amino acids is mediated by the system y+ family and the cationic amino acid transporter (CAT) family. Members of the CAT family share a high degree of sequence homology, and each contains 12-14 putative transmembrane domains. (Ito, K. and Groudine, M. (1997) J. Biol. Chem. 272:26780-26786.)

Hormones are secreted molecules that travel through the circulation and bind to specific receptors on the surface of, or within, target cells. Although they have diverse biochemical compositions

neurotrophins, growth factors, interferons, and chemokines all define cytokine families that work in conjunction with cellular receptors to regulate cell proliferation and differentiation. In addition, cytokines effect activities such as leukocyte migration and function, hematopoietic cell proliferation, temperature regulation, acute response to infection, tissue remodeling, and apoptosis.

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Chemokines, in particular, are small chemoattractant cytokines involved in inflammation, leukocyte proliferation and migration, angiogenesis and angiostasis, regulation of hematopoiesis, HIV infectivity, and stimulation of cytokine secretion. Chemokines generally contain 70-100 amino acids and are subdivided into four subfamilies based on the presence of conserved cysteine-based motifs. (Callard, R. and Gearing, A. (1994) The Cytokine Facts Book, Academic Press, New York, NY, pp. 181-190, 210-213, 223-227.)

Growth and differentiation factors are secreted proteins which function in intercellular communication. Some factors require oligomerization or association with MPs for activity. Complex interactions among these factors and their receptors trigger intracellular signal transduction pathways that stimulate or inhibit cell division, cell differentiation, cell signaling, and cell motility. Most growth and differentiation factors act on cells in their local environment (paracrine signaling). There are three broad classes of growth and differentiation factors. The first class includes the large polypeptide growth factors such as epidermal growth factor, fibroblast growth factor, transforming growth factor, insulin-like growth factor, and platelet-derived growth factor. The second class includes the hematopoietic growth factors such as the colony stimulating factors (CSFs). Hematopoietic growth factors stimulate the proliferation and differentiation of blood cells such as B-lymphocytes, T-lymphocytes, erythrocytes, platelets, eosinophils, basophils, neutrophils, macrophages, and their stem cell precursors. The third class includes small peptide factors such as bombesin, vasopressin, oxytocin, endothelin, transferrin, angiotensin II, vasoactive intestinal peptide, and bradykinin which function as hormones to regulate cellular functions other than proliferation.

Growth and differentiation factors play critical roles in neoplastic transformation of cells in vitro and in tumor progression in vivo. Inappropriate expression of growth factors by tumor cells may contribute to vascularization and metastasis of tumors. During hematopoiesis, growth factor misregulation can result in anemias, leukemias, and lymphomas. Certain growth factors such as interferon are cytotoxic to tumor cells both in vivo and in vitro. Moreover, some growth factors and growth factor receptors are related both structurally and functionally to oncoproteins. In addition, growth factors affect transcriptional regulation of both proto-oncogenes and oncosuppressor genes. (Reviewed in Pimentel, E. (1994) Handbook of Growth Factors, CRC Press, Ann Arbor, MI, pp. 1-9.)

Proteolytic enzymes or proteases either activate or deactivate proteins by hydrolyzing peptide bonds. Proteases are found in the cytosol, in membrane-bound compartments, and in the extracellular space. The major families are the zinc, serine, cysteine, thiol, and carboxyl proteases.

e) an RNA equivalent of a) through d). In one alternative, the polynucleotide comprises a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63. In another alternative, the polynucleotide comprises at least 60 contiguous nucleotides of a polynucleotide sequence selected from the group consisting of a) a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; b) a naturally occurring polynucleotide sequence having at least 90% sequence identity to a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; c) a polynucleotide sequence complementary to a); d) a polynucleotide sequence complementary to b); and e) an RNA equivalent of a) through d). The invention further provides a composition for the detection of expression of secretory polynucleotides comprising at least one isolated polynucleotide comprising a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; b) a naturally occurring polynucleotide sequence having at least 90% sequence identity to a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; c) a polynucleotide sequence complementary to a); d) a polynucleotide sequence complementary to b); and e) an RNA equivalent of a) through d); and a detectable label.

The invention also provides a method for detecting a target polynucleotide in a sample, said target polynucleotide comprising a polynucleotide sequence selected from the group consisting of a) a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; b) a naturally occurring polynucleotide sequence having at least 90% sequence identity to a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; c) a polynucleotide sequence complementary to a); d) a polynucleotide sequence complementary to b); and e) an RNA equivalent of a) through d). The method comprises a) hybridizing the sample with a probe comprising at least 20 contiguous nucleotides comprising a sequence complementary to said target polynucleotide in the sample, and which probe specifically hybridizes to said target polynucleotide, under conditions whereby a hybridization complex is formed between said probe and said target polynucleotide, and b) detecting the presence or absence of said hybridization complex, and, optionally, if present, the amount thereof. In one alternative, the probe comprises at least 30 contiguous nucleotides. In another alternative, the probe comprises at least 60 contiguous nucleotides.

The invention further provides a recombinant polynucleotide comprising a promoter sequence operably linked to an isolated polynucleotide comprising a polynucleotide sequence selected from the group consisting of a) a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; b) a naturally occurring polynucleotide sequence having at least 90% sequence identity to a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; c) a polynucleotide sequence complementary to a); d) a polynucleotide sequence complementary to b); and e) an RNA equivalent of a) through d). In one alternative, the invention provides a cell transformed with the recombinant polynucleotide. In another alternative, the invention provides a transgenic organism

and mechanisms of action, hormones can be grouped into two categories. One category includes small lipophilic hormones that diffuse through the plasma membrane of target cells, bind to cytosolic or nuclear receptors, and form a complex that alters gene expression. Examples of these molecules include retinoic acid, thyroxine, and the cholesterol-derived steroid hormones such as progesterone, estrogen, testosterone, cortisol, and aldosterone. The second category includes hydrophilic hormones that function by binding to cell surface receptors that transduce signals across the plasma membrane. Examples of such hormones include amino acid derivatives such as catecholamines and peptide hormones such as glucagon, insulin, gastrin, secretin, cholecystokinin, adrenocorticotropic hormone, follicle stimulating hormone, luteinizing hormone, thyroid stimulating hormone, and vasopressin. (See, for example, Lodish et al. (1995) Molecular Cell Biology, Scientific American Books Inc., New York, NY, pp. 856-864.)

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Neuropeptides and vasomediators (NP/VM) comprise a large family of endogenous signaling molecules. Included in this family are neuropeptides and neuropeptide hormones such as bombesin, neuropeptide Y, neurotensin, neuromedin N, melanocortins, opioids, galanin, somatostatin, tachykinins, urotensin II and related peptides involved in smooth muscle stimulation, vasopressin, vasoactive intestinal peptide, and circulatory system-borne signaling molecules such as angiotensin, complement, calcitonin, endothelins, formyl-methionyl peptides, glucagon, cholecystokinin and gastrin. NP/VMs can transduce signals directly, modulate the activity or release of other neurotransmitters and hormones, and act as catalytic enzymes in cascades. The effects of NP/VMs range from extremely brief to long-lasting. (Reviewed in Martin, C. R. et al. (1985) Endocrine Physiology, Oxford University Press, New York, NY, pp. 57-62.)

The discovery of new secretory molecules satisfies a need in the art by providing new compositions which are useful in the diagnosis, study, prevention, and treatment of diseases associated with, as well as effects of exogenous compounds on, cell signaling and the expression of secretory molecules.

SUMMARY OF THE INVENTION

The present invention relates to nucleic acid sequences comprising human polynucleotides encoding secretory polypeptides that contain signal peptides and/or transmembrane domains. These human polynucleotides (sptm) as presented in the Sequence Listing uniquely identify partial or full length genes encoding structural, functional, and regulatory polypeptides involved in cell signaling.

The invention provides an isolated polynucleotide comprising a polynucleotide sequence selected from the group consisting of a) a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; b) a naturally occurring polynucleotide sequence having at least 90% sequence identity to a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; c) a polynucleotide sequence complementary to a); d) a polynucleotide sequence complementary to b); and

contiguous nucleotides of a polynucleotide comprising a polynucleotide sequence selected from the group consisting of i) a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; ii) a naturally occurring polynucleotide sequence having at least 90% sequence identity to a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; iii) a polynucleotide sequence complementary to i), iv) a polynucleotide sequence complementary to ii), and v) an RNA equivalent of i)-iv). Hybridization occurs under conditions whereby a specific hybridization complex is formed between said probe and a target polynucleotide in the biological sample, said target polynucleotide comprising a polynucleotide sequence selected from the group consisting of i) a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; ii) a naturally occurring polynucleotide sequence having at least 90% sequence identity to a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; iii) a polynucleotide sequence complementary to i), iv) a polynucleotide sequence complementary to ii), and v) an RNA equivalent of i)-iv), and alternatively, the target polynucleotide comprises a fragment of a polynucleotide sequence selected from the group consisting of i-v above; c) quantifying the amount of hybridization complex; and d) comparing the amount of hybridization complex in the treated biological sample with the amount of hybridization complex in an untreated biological sample, wherein a difference in the amount of hybridization complex in the treated biological sample is indicative of toxicity of the test compound.

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DESCRIPTION OF THE TABLES

Table 1 shows the sequence identification numbers (SEQ ID NO:s) and template identification numbers (template IDs) corresponding to the polynucleotides of the present invention, along with polynucleotide segments of each template sequence as defined by the indicated "start" and "stop" nucleotide positions. The reading frames of the polynucleotide segments are shown, and the polypeptides encoded by the polynucleotide segments constitute either signal peptide (SP) or transmembrane (TM) domains, as indicated.

Table 2 shows the sequence identification numbers (SEQ ID NO:s) and template identification numbers (template IDs) corresponding to the polynucleotides of the present invention, along with component sequence identification numbers (component IDs) corresponding to each template. The component sequences, which were used to assemble the template sequences, are defined by the indicated "start" and "stop" nucleotide positions along each template.

Table 3 shows the tissue distribution profiles for the templates of the invention.

Table 4 summarizes the bioinformatics tools which are useful for analysis of the polynucleotides of the present invention. The first column of Table 4 lists analytical tools, programs, and algorithms, the second column provides brief descriptions thereof, the third column presents appropriate references, all of which are incorporated by reference herein in their entirety, and the fourth column presents, where applicable, the scores, probability values, and other parameters used to evaluate

comprising the recombinant polynucleotide. In a further alternative, the invention provides a method for producing a secretory polypeptide, the method comprising a) culturing a cell under conditions suitable for expression of the secretory polypeptide, wherein said cell is transformed with the recombinant polynucleotide, and b) recovering the secretory polypeptide so expressed.

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The invention also provides a purified secretory polypeptide (SPTM) encoded by at least one polynucleotide comprising a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63. Additionally, the invention provides an isolated antibody which specifically binds to the secretory polypeptide. The invention further provides a method of identifying a test compound which specifically binds to the secretory polypeptide, the method comprising the steps of a) providing a test compound; b) combining the secretory polypeptide with the test compound for a sufficient time and under suitable conditions for binding; and c) detecting binding of the secretory polypeptide to the test compound, thereby identifying the test compound which specifically binds the secretory polypeptide.

The invention further provides a microarray wherein at least one element of the microarray is an isolated polynucleotide comprising at least 60 contiguous nucleotides of a polynucleotide comprising a polynucleotide sequence selected from the group consisting of a) a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; b) a naturally occurring polynucleotide sequence having at least 90% sequence identity to a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; c) a polynucleotide sequence complementary to a); d) a polynucleotide sequence complementary to b); and e) an RNA equivalent of a) through d). The invention also provides a method for generating a transcript image of a sample which contains polynucleotides. The method comprises a) labeling the polynucleotides of the sample, b) contacting the elements of the microarray with the labeled polynucleotides of the sample under conditions suitable for the formation of a hybridization complex, and c) quantifying the expression of the polynucleotides in the sample.

Additionally, the invention provides a method for screening a compound for effectiveness in altering expression of a target polynucleotide, wherein said target polynucleotide comprises a polynucleotide sequence selected from the group consisting of a) a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; b) a naturally occurring polynucleotide sequence having at least 90% sequence identity to a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63; c) a polynucleotide sequence complementary to a); d) a polynucleotide sequence complementary to b); and e) an RNA equivalent of a) through d). The method comprises a) exposing a sample comprising the target polynucleotide to a compound, and b) detecting altered expression of the target polynucleotide.

The invention further provides a method for assessing toxicity of a test compound, said method comprising a) treating a biological sample containing nucleic acids with the test compound; b) hybridizing the nucleic acids of the treated biological sample with a probe comprising at least 20

Fv fragments, which are capable of binding the epitopic determinant. Antibodies that bind SPTM polypeptides can be prepared using intact polypeptides or using fragments containing small peptides of interest as the immunizing antigen. The polypeptide or peptide used to immunize an animal (e.g., a mouse, a rat, or a rabbit) can be derived from the translation of RNA, or synthesized chemically, and can be conjugated to a carrier protein if desired. Commonly used carriers that are chemically coupled to peptides include bovine serum albumin, thyroglobulin, and keyhole limpet hemocyanin (KLH). The coupled peptide is then used to immunize the animal.

"Antisense sequence" refers to a sequence capable of specifically hybridizing to a target sequence. The antisense sequence may include DNA, RNA, or any nucleic acid mimic or analog such as peptide nucleic acid (PNA); oligonucleotides having modified backbone linkages such as phosphorothioates, methylphosphonates, or benzylphosphonates; oligonucleotides having modified sugar groups such as 2'-methoxyethyl sugars or 2'-methoxyethoxy sugars; or oligonucleotides having modified bases such as 5-methyl cytosine, 2'-deoxyuracil, or 7-deaza-2'-deoxyguanosine.

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"Antisense sequence" refers to a sequence capable of specifically hybridizing to a target sequence. The antisense sequence can be DNA, RNA, or any nucleic acid mimic or analog.

"Antisense technology" refers to any technology which relies on the specific hybridization of an antisense sequence to a target sequence.

A "bin" is a portion of computer memory space used by a computer program for storage of data, and bounded in such a manner that data stored in a bin may be retrieved by the program.

"Biologically active" refers to an amino acid sequence having a structural, regulatory, or biochemical function of a naturally occurring amino acid sequence.

"Clone joining" is a process for combining gene bins based upon the bins' containing sequence information from the same clone. The sequences may assemble into a primary gene transcript as well as one or more splice variants.

"Complementary" describes the relationship between two single-stranded nucleic acid sequences that anneal by base-pairing (5'-A-G-T-3' pairs with its complement 3'-T-C-A-5').

A "component sequence" is a nucleic acid sequence selected by a computer program such as PHRED and used to assemble a consensus or template sequence from one or more component sequences.

A "consensus sequence" or "template sequence" is a nucleic acid sequence which has been assembled from overlapping sequences, using a computer program for fragment assembly such as the GELVIEW fragment assembly system (Genetics Computer Group (GCG), Madison WI) or using a relational database management system (RDMS).

"Conservative amino acid substitutions" are those substitutions that, when made, least interfere with the properties of the original protein, i.e., the structure and especially the function of the protein is

the strength of a match between two sequences (the higher the score, the greater the homology between

DETAILED DESCRIPTION OF THE INVENTION

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Before the nucleic acid sequences and methods are presented, it is to be understood that this invention is not limited to the particular machines, methods, and materials described. Although particular embodiments are described, machines, methods, and materials similar or equivalent to these embodiments may be used to practice the invention. The preferred machines, methods, and materials set forth are not intended to limit the scope of the invention which is limited only by the appended claims.

The singular forms "a", "an", and "the" include plural reference unless the context clearly 10 dictates otherwise. All technical and scientific terms have the meanings commonly understood by one of ordinary skill in the art. All publications are incorporated by reference for the purpose of describing and disclosing the cell lines, vectors, and methodologies which are presented and which might be used in connection with the invention. Nothing in the specification is to be construed as an admission that the invention is not entitled to antedate such disclosure by virtue of prior invention. 15 **Definitions**

As used herein, the lower case "sptm" refers to a nucleic acid sequence, while the upper case "SPTM" refers to an amino acid sequence encoded by sptm. A "full-length" sptm refers to a nucleic acid sequence containing the entire coding region of a gene endogenously expressed in human tissue.

"Adjuvants" are materials such as Freund's adjuvant, mineral gels (aluminum hydroxide), and surface active substances (lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, keyhole limpet hemocyanin, and dinitrophenol) which may be administered to increase a host's immunological response.

"Allele" refers to an alternative form of a nucleic acid sequence. Alleles result from a "mutation," a change or an alternative reading of the genetic code. Any given gene may have none, one, 25 or many allelic forms. Mutations which give rise to alleles include deletions, additions, or substitutions of nucleotides. Each of these changes may occur alone, or in combination with the others, one or more times in a given nucleic acid sequence. The present invention encompasses allelic sptm.

"Amino acid sequence" refers to a peptide, a polypeptide, or a protein of either natural or synthetic origin. The amino acid sequence is not limited to the complete, endogenous amino acid sequence and may be a fragment, epitope, variant, or derivative of a protein expressed by a nucleic acid sequence.

"Amplification" refers to the production of additional copies of a sequence and is carried out using polymerase chain reaction (PCR) technologies well known in the art.

"Antibody" refers to intact molecules as well as to fragments thereof, such as Fab, F(ab')2, and 35

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be preferentially selected from certain regions of a molecule. For example, a polypeptide fragment may comprise a certain length of contiguous amino acids selected from the first 250 or 500 amino acids (or first 25% or 50%) of a polypeptide as shown in a certain defined sequence. Clearly these lengths are exemplary, and any length that is supported by the specification, including the Sequence Listing and the figures, may be encompassed by the present embodiments.

A fragment of sptm comprises a region of unique polynucleotide sequence that specifically identifies sptm, for example, as distinct from any other sequence in the same genome. A fragment of sptm is useful, for example, in hybridization and amplification technologies and in analogous methods that distinguish sptm from related polynucleotide sequences. The precise length of a fragment of sptm and the region of sptm to which the fragment corresponds are routinely determinable by one of ordinary skill in the art based on the intended purpose for the fragment.

A fragment of SPTM is encoded by a fragment of sptm. A fragment of SPTM comprises a region of unique amino acid sequence that specifically identifies SPTM. For example, a fragment of SPTM is useful as an immunogenic peptide for the development of antibodies that specifically recognize SPTM. The precise length of a fragment of SPTM and the region of SPTM to which the fragment corresponds are routinely determinable by one of ordinary skill in the art based on the intended purpose for the fragment.

A "full length" nucleotide sequence is one containing at least a start site for translation to a protein sequence, followed by an open reading frame and a stop site, and encoding a "full length" polypeptide.

"Hit" refers to a sequence whose annotation will be used to describe a given template. Criteria for selecting the top hit are as follows: if the template has one or more exact nucleic acid matches, the top hit is the exact match with highest percent identity. If the template has no exact matches but has significant protein hits, the top hit is the protein hit with the lowest E-value. If the template has no significant protein hits, but does have significant non-exact nucleotide hits, the top hit is the nucleotide hit with the lowest E-value.

"Homology" refers to sequence similarity either between a reference nucleic acid sequence and at least a fragment of an sptm or between a reference amino acid sequence and a fragment of an SPTM.

"Hybridization" refers to the process by which a strand of nucleotides anneals with a complementary strand through base pairing. Specific hybridization is an indication that two nucleic acid sequences share a high degree of identity. Specific hybridization complexes form under defined annealing conditions, and remain hybridized after the "washing" step. The defined hybridization conditions include the annealing conditions and the washing step(s), the latter of which is particularly important in determining the stringency of the hybridization process, with more stringent conditions allowing less non-specific binding, i.e., binding between pairs of nucleic acid probes that are not

conserved and not significantly changed by such substitutions. The table below shows amino acids which may be substituted for an original amino acid in a protein and which are regarded as conservative substitutions

5	Original Residue	Comme
	Ala	Conservative Substitution
	Arg ·	Gly, Ser
	Asn	His, Lys
10	Asp	Asp, Gln, His
	Cys	Asn, Glu
	Gln	Ala, Ser
	Glu	Asn, Glu, His
	Gly	Asp, Gln, His
	His	Ala
15	Пе	Asn, Arg, Gln, Glu
	Leu	Leu, Val
	Lys	Ile, Val
	Met	Arg, Gln, Glu
	Phe	Leu, Ile
20	Ser	His, Met, Leu, Trp, Tyr
	Thr	Cys, Thr
	T _{TP}	Ser, Val
	Tyr	Phe, Tyr
	Val	His, Phe, Trp
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	Conservative substitutions generally magnetic	aintain (a) the structure of the

Conservative substitutions generally maintain (a) the structure of the polypeptide backbone in the area of the substitution, for example, as a beta sheet or alpha helical conformation, (b) the charge or hydrophobicity of the molecule at the target site, or (c) the bulk of the side chain.

"Deletion" refers to a change in either a nucleic or amino acid sequence in which at least one nucleotide or amino acid residue, respectively, is absent.

"Derivative" refers to the chemical modification of a nucleic acid sequence, such as by replacement of hydrogen by an alkyl, acyl, amino, hydroxyl, or other group.

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The terms "element" and "array element" refer to a polynucleotide, polypeptide, or other chemical compound having a unique and defined position on a microarray.

"E-value" refers to the statistical probability that a match between two sequences occurred by

A "fragment" is a unique portion of sptm or SPTM which is identical in sequence to but shorter in length than the parent sequence. A fragment may comprise up to the entire length of the defined sequence, minus one nucleotide/amino acid residue. For example, a fragment may comprise from 10 to 1000 contiguous amino acid residues or nucleotides. A fragment used as a probe, primer, antigen, therapeutic molecule, or for other purposes, may be at least 5, 10, 15, 16, 20, 25, 30, 40, 50, 60, 75, 100, 150, 250 or at least 500 contiguous amino acid residues or nucleotides in length. Fragments may

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multiple restriction enzyme sites and to provide for the use of enzymes which leave 5' or 3' overhangs (e.g., BamHI, EcoRI, and HindIII) and those which provide blunt ends (e.g., EcoRV, SnaBI, and Stul).

"Naturally occurring" refers to an endogenous polynucleotide or polypeptide that may be isolated from viruses or prokaryotic or eukaryotic cells.

"Nucleic acid sequence" refers to the specific order of nucleotides joined by phosphodiester bonds in a linear, polymeric arrangement. Depending on the number of nucleotides, the nucleic acid sequence can be considered an oligomer, oligonucleotide, or polynucleotide. The nucleic acid can be DNA, RNA, or any nucleic acid analog, such as PNA, may be of genomic or synthetic origin, may be either double-stranded or single-stranded, and can represent either the sense or antisense (complementary) strand.

"Oligomer" refers to a nucleic acid sequence of at least about 6 nucleotides and as many as about 60 nucleotides, preferably about 15 to 40 nucleotides, and most preferably between about 20 and 30 nucleotides, that may be used in hybridization or amplification technologies. Oligomers may be used as, e.g., primers for PCR, and are usually chemically synthesized.

"Operably linked" refers to the situation in which a first nucleic acid sequence is placed in a functional relationship with the second nucleic acid sequence. For instance, a promoter is operably linked to a coding sequence if the promoter affects the transcription or expression of the coding sequence. Generally, operably linked DNA sequences may be in close proximity or contiguous and, where necessary to join two protein coding regions, in the same reading frame.

"Peptide nucleic acid" (PNA) refers to a DNA mimic in which nucleotide bases are attached to a pseudopeptide backbone to increase stability. PNAs, also designated antigene agents, can prevent gene expression by targeting complementary messenger RNA.

The phrases "percent identity" and "% identity", as applied to polynucleotide sequences, refer to the percentage of residue matches between at least two polynucleotide sequences aligned using a standardized algorithm. Such an algorithm may insert, in a standardized and reproducible way, gaps in the sequences being compared in order to optimize alignment between two sequences, and therefore achieve a more meaningful comparison of the two sequences.

Percent identity between polynucleotide sequences may be determined using the default parameters of the CLUSTAL V algorithm as incorporated into the MEGALIGN version 3.12e sequence alignment program. This program is part of the LASERGENE software package, a suite of molecular biological analysis programs (DNASTAR, Madison WI). CLUSTAL V is described in Higgins, D.G. and Sharp, P.M. (1989) CABIOS 5:151-153 and in Higgins, D.G. et al. (1992) CABIOS 8:189-191. For pairwise alignments of polynucleotide sequences, the default parameters are set as follows: Ktuple=2, gap penalty=5, window=4, and "diagonals saved"=4. The "weighted" residue weight table is

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perfectly matched. Permissive conditions for annealing of nucleic acid sequences are routinely determinable and may be consistent among hybridization experiments, whereas wash conditions may be varied among experiments to achieve the desired stringency.

Generally, stringency of hybridization is expressed with reference to the temperature under which the wash step is carried out. Generally, such wash temperatures are selected to be about 5°C to 20°C lower than the thermal melting point (T_m) for the specific sequence at a defined ionic strength and pH. The T_m is the temperature (under defined ionic strength and pH) at which 50% of the target sequence hybridizes to a perfectly matched probe. An equation for calculating T_m and conditions for nucleic acid hybridization is well known and can be found in Sambrook et al., 1989, Molecular Cloning: A Laboratory Manual, 2nd ed., vol. 1-3, Cold Spring Harbor Press, Plainview NY; specifically see volume 2, chapter 9.

High stringency conditions for hybridization between polynucleotides of the present invention include wash conditions of 68° C in the presence of about $0.2 \times SSC$ and about 0.1% SDS, for 1 hour. Alternatively, temperatures of about 65° C, 60° C, or 55° C may be used. SSC concentration may be varied from about 0.2 to $2 \times SSC$, with SDS being present at about 0.1%. Typically, blocking reagents are used to block non-specific hybridization. Such blocking reagents include, for instance, denatured salmon sperm DNA at about $100\text{-}200~\mu\text{g/ml}$. Useful variations on these conditions will be readily apparent to those skilled in the art. Hybridization, particularly under high stringency conditions, may be suggestive of evolutionary similarity between the nucleotides. Such similarity is strongly indicative of a similar role for the nucleotides and their resultant proteins.

Other parameters, such as temperature, salt concentration, and detergent concentration may be varied to achieve the desired stringency. Denaturants, such as formamide at a concentration of about 35-50% v/v, may also be used under particular circumstances, such as RNA:DNA hybridizations. Appropriate hybridization conditions are routinely determinable by one of ordinary skill in the art.

"Immunogenic" describes the potential for a natural, recombinant, or synthetic peptide, epitope, polypeptide, or protein to induce antibody production in appropriate animals, cells, or cell lines.

"Insertion" or "addition" refers to a change in either a nucleic or amino acid sequence in which at least one nucleotide or residue, respectively, is added to the sequence.

"Labeling" refers to the covalent or noncovalent joining of a polynucleotide, polypeptide, or antibody with a reporter molecule capable of producing a detectable or measurable signal.

"Microarray" is any arrangement of nucleic acids, amino acids, antibodies, etc., on a substrate. The substrate may be a solid support such as beads, glass, paper, nitrocellulose, nylon, or an appropriate membrane.

"Linkers" are short stretches of nucleotide sequence which may be added to a vector or an sptm to create restriction endonuclease sites to facilitate cloning. "Polylinkers" are engineered to incorporate

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The phrases "percent identity" and "% identity", as applied to polypeptide sequences, refer to the percentage of residue matches between at least two polypeptide sequences aligned using a standardized algorithm. Methods of polypeptide sequence alignment are well-known. Some alignment methods take into account conservative amino acid substitutions. Such conservative substitutions, explained in more detail above, generally preserve the hydrophobicity and acidity of the substituted residue, thus preserving the structure (and therefore function) of the folded polypeptide.

Percent identity between polypeptide sequences may be determined using the default parameters of the CLUSTAL V algorithm as incorporated into the MEGALIGN version 3.12e sequence alignment program (described and referenced above). For pairwise alignments of polypeptide sequences using CLUSTAL V, the default parameters are set as follows: Ktuple=1, gap penalty=3, window=5, and "diagonals saved"=5. The PAM250 matrix is selected as the default residue weight table. As with polynucleotide alignments, the percent identity is reported by CLUSTAL V as the "percent similarity" between aligned polypeptide sequence pairs.

Alternatively the NCBI BLAST software suite may be used. For example, for a pairwise comparison of two polypeptide sequences, one may use the "BLAST 2 Sequences" tool Version 2.0.9 (May-07-1999) with blastp set at default parameters. Such default parameters may be, for example:

Matrix: BLOSUM62

Open Gap: 11 and Extension Gap: 1 penalty

Gap x drop-off: 50

20 Expect: 10

Word Size: 3

Filter: on

Percent identity may be measured over the length of an entire defined polypeptide sequence, for example, as defined by a particular SEQ ID number, or may be measured over a shorter length, for example, over the length of a fragment taken from a larger, defined polypeptide sequence, for instance, a fragment of at least 15, at least 20, at least 30, at least 40, at least 50, at least 70 or at least 150 contiguous residues. Such lengths are exemplary only, and it is understood that any fragment length supported by the sequences shown herein, in figures or Sequence Listings, may be used to describe a length over which percentage identity may be measured.

"Post-translational modification" of an SPTM may involve lipidation, glycosylation, phosphorylation, acetylation, racemization, proteolytic cleavage, and other modifications known in the art. These processes may occur synthetically or biochemically. Biochemical modifications will vary by cell type depending on the enzymatic milieu and the SPTM.

"Probe" refers to sptm or fragments thereof, which are used to detect identical, allelic or related nucleic acid sequences. Probes are isolated oligonucleotides or polynucleotides attached to a detectable

selected as the default. Percent identity is reported by CLUSTAL V as the "percent similarity" between aligned polynucleotide sequence pairs.

Alternatively, a suite of commonly used and freely available sequence comparison algorithms is provided by the National Center for Biotechnology Information (NCBI) Basic Local Alignment Search Tool (BLAST) (Altschul, S.F. et al. (1990) J. Mol. Biol. 215:403-410), which is available from several sources, including the NCBI, Bethesda, MD, and on the Internet at http://www.ncbi.nlm.nih.gov/BLAST/. The BLAST software suite includes various sequence analysis programs including "blastn," that is used to determine alignment between a known polynucleotide sequence and other sequences on a variety of databases. Also available is a tool called "BLAST 2 Sequences" that is used for direct pairwise comparison of two nucleotide sequences. "BLAST 2 10 Sequences" can be accessed and used interactively at http://www.ncbi.nlm.nih.gov/gorf/bl2/. The "BLAST 2 Sequences" tool can be used for both blastn and blastp (discussed below). BLAST programs are commonly used with gap and other parameters set to default settings. For example, to compare two nucleotide sequences, one may use blastn with the "BLAST 2 Sequences" tool Version 2.0.9 (May-07-1999) set at default parameters. Such default parameters may be, for example: 15

Matrix: BLOSUM62

Reward for match: 1

Penalty for mismatch: -2

Open Gap: 5 and Extension Gap: 2 penalties

20 Gap x drop-off: 50

Expect: 10

Word Size: 11

Filter: on

Percent identity may be measured over the length of an entire defined sequence, for example, as defined by a particular SEQ ID number, or may be measured over a shorter length, for example, over 25 the length of a fragment taken from a larger, defined sequence, for instance, a fragment of at least 20, at least 30, at least 40, at least 50, at least 70, at least 100, or at least 200 contiguous nucleotides. Such lengths are exemplary only, and it is understood that any fragment length supported by the sequences shown herein, in figures or Sequence Listings, may be used to describe a length over which percentage identity may be measured.

Nucleic acid sequences that do not show a high degree of identity may nevertheless encode similar amino acid sequences due to the degeneracy of the genetic code. It is understood that changes in nucleic acid sequence can be made using this degeneracy to produce multiple nucleic acid sequences that all encode substantially the same protein.

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polynucleotide fragments. The oligonucleotides and polynucleotide fragments identified by any of the above selection methods are useful in hybridization technologies, for example, as PCR or sequencing primers, microarray elements, or specific probes to identify fully or partially complementary polynucleotides in a sample of nucleic acids. Methods of oligonucleotide selection are not limited to those described above.

"Purified" refers to molecules, either polynucleotides or polypeptides that are isolated or separated from their natural environment and are at least 60% free, preferably at least 75% free, and most preferably at least 90% free from other compounds with which they are naturally associated.

A "recombinant nucleic acid" is a sequence that is not naturally occurring or has a sequence that is made by an artificial combination of two or more otherwise separated segments of sequence. This artificial combination is often accomplished by chemical synthesis or, more commonly, by the artificial manipulation of isolated segments of nucleic acids, e.g., by genetic engineering techniques such as those described in Sambrook, <u>supra</u>. The term recombinant includes nucleic acids that have been altered solely by addition, substitution, or deletion of a portion of the nucleic acid. Frequently, a recombinant nucleic acid may include a nucleic acid sequence operably linked to a promoter sequence. Such a recombinant nucleic acid may be part of a vector that is used, for example, to transform a cell.

Alternatively, such recombinant nucleic acids may be part of a viral vector, e.g., based on a vaccinia virus, that could be use to vaccinate a mammal wherein the recombinant nucleic acid is expressed, inducing a protective immunological response in the mammal.

"Regulatory element" refers to a nucleic acid sequence from nontranslated regions of a gene, and includes enhancers, promoters, introns, and 3' untranslated regions, which interact with host proteins to carry out or regulate transcription or translation.

"Reporter" molecules are chemical or biochemical moieties used for labeling a nucleic acid, an amino acid, or an antibody. They include radionuclides; enzymes; fluorescent, chemiluminescent, or chromogenic agents; substrates; cofactors; inhibitors; magnetic particles; and other moieties known in the art.

An "RNA equivalent," in reference to a DNA sequence, is composed of the same linear sequence of nucleotides as the reference DNA sequence with the exception that all occurrences of the nitrogenous base thymine are replaced with uracil, and the sugar backbone is composed of ribose instead of deoxyribose.

"Sample" is used in its broadest sense. Samples may contain nucleic or amino acids, antibodies, or other materials, and may be derived from any source (e.g., bodily fluids including, but not limited to, saliva, blood, and urine; chromosome(s), organelles, or membranes isolated from a cell; genomic DNA, RNA, or cDNA in solution or bound to a substrate; and cleared cells or tissues or blots or imprints from such cells or tissues).

label or reporter molecule. Typical labels include radioactive isotopes, ligands, chemiluminescent agents, and enzymes. "Primers" are short nucleic acids, usually DNA oligonucleotides, which may be annealed to a target polynucleotide by complementary base-pairing. The primer may then be extended along the target DNA strand by a DNA polymerase enzyme. Primer pairs can be used for amplification (and identification) of a nucleic acid sequence, e.g., by the polymerase chain reaction (PCR).

Probes and primers as used in the present invention typically comprise at least 15 contiguous nucleotides of a known sequence. In order to enhance specificity, longer probes and primers may also be employed, such as probes and primers that comprise at least 20, 30, 40, 50, 60, 70, 80, 90, 100, or at least 150 consecutive nucleotides of the disclosed nucleic acid sequences. Probes and primers may be considerably longer than these examples, and it is understood that any length supported by the specification, including the figures and Sequence Listing, may be used.

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Methods for preparing and using probes and primers are described in the references, for example Sambrook et al., 1989, Molecular Cloning: A Laboratory Manual, 2nd ed., vol. 1-3, Cold Spring Harbor Press, Plainview NY; Ausubel et al., 1987, Current Protocols in Molecular Biology,

Greene Publ. Assoc. & Wiley-Intersciences, New York NY; Innis et al., 1990, PCR Protocols, A Guide to Methods and Applications, Academic Press, San Diego CA. PCR primer pairs can be derived from a known sequence, for example, by using computer programs intended for that purpose such as Primer (Version 0.5, 1991, Whitehead Institute for Biomedical Research, Cambridge MA).

Oligonucleotides for use as primers are selected using software known in the art for such purpose. For example, OLIGO 4.06 software is useful for the selection of PCR primer pairs of up to 20 100 nucleotides each, and for the analysis of oligonucleotides and larger polynucleotides of up to 5,000 nucleotides from an input polynucleotide sequence of up to 32 kilobases. Similar primer selection programs have incorporated additional features for expanded capabilities. For example, the PrimOU primer selection program (available to the public from the Genome Center at University of Texas South West Medical Center, Dallas TX) is capable of choosing specific primers from megabase sequences 25 and is thus useful for designing primers on a genome-wide scope. The Primer3 primer selection program (available to the public from the Whitehead Institute/MIT Center for Genome Research, Cambridge MA) allows the user to input a "mispriming library," in which sequences to avoid as primer binding sites are user-specified. Primer3 is useful, in particular, for the selection of oligonucleotides for microarrays. (The source code for the latter two primer selection programs may also be obtained from 30 their respective sources and modified to meet the user's specific needs.) The PrimeGen program (available to the public from the UK Human Genome Mapping Project Resource Centre, Cambridge UK) designs primers based on multiple sequence alignments, thereby allowing selection of primers that hybridize to either the most conserved or least conserved regions of aligned nucleic acid sequences. Hence, this program is useful for identification of both unique and conserved oligonucleotides and 35

A "variant" of a particular nucleic acid sequence is defined as a nucleic acid sequence having at least 25% sequence identity to the particular nucleic acid sequence over a certain length of one of the nucleic acid sequences using blastn with the "BLAST 2 Sequences" tool Version 2.0.9 (May-07-1999) set at default parameters. Such a pair of nucleic acids may show, for example, at least 30%, at least 50%, at least 60%, at least 70%, at least 80%, at least 95% or even at least 98% or greater sequence identity over a certain defined length. The variant may result in "conservative" amino acid changes which do not affect structural and/or chemical properties. A variant may be described as, for example, an "allelic" (as defined above), "splice," "species," or "polymorphic" variant. A splice variant may have significant identity to a reference molecule, but will generally have a greater or lesser number of polynucleotides due to alternate splicing of exons during mRNA processing. The corresponding polypeptide may possess additional functional domains or lack domains that are present in the reference molecule. Species variants are polynucleotide sequences that vary from one species to another. The resulting polypeptides generally will have significant amino acid identity relative to each other. A polymorphic variant is a variation in the polymucleotide sequence of a particular gene between individuals of a given species. Polymorphic variants also may encompass "single nucleotide polymorphisms" (SNPs) in which the polynucleotide sequence varies by one base. The presence of SNPs may be indicative of, for example, a certain population, a disease state, or a propensity for a disease state.

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In an alternative, variants of the polynucleotides of the present invention may be generated through recombinant methods. One possible method is a DNA shuffling technique such as MOLECULARBREEDING (Maxygen Inc., Santa Clara CA; described in U.S. Patent Number 5,837,458; Chang, C.-C. et al. (1999) Nat. Biotechnol. 17:793-797; Christians, F.C. et al. (1999) Nat. Biotechnol. 17:259-264; and Crameri, A. et al. (1996) Nat. Biotechnol. 14:315-319) to alter or improve the biological properties of SPTM, such as its biological or enzymatic activity or its ability to bind to other molecules or compounds. DNA shuffling is a process by which a library of gene variants is 25 produced using PCR-mediated recombination of gene fragments. The library is then subjected to selection or screening procedures that identify those gene variants with the desired properties. These preferred variants may then be pooled and further subjected to recursive rounds of DNA shuffling and selection/screening. Thus, genetic diversity is created through "artificial" breeding and rapid molecular evolution. For example, fragments of a single gene containing random point mutations may be recombined, screened, and then reshuffled until the desired properties are optimized. Alternatively, fragments of a given gene may be recombined with fragments of homologous genes in the same gene family, either from the same or different species, thereby maximizing the genetic diversity of multiple naturally occurring genes in a directed and controllable manner.

"Specific binding" or "specifically binding" refers to the interaction between a protein or peptide and its agonist, antibody, antagonist, or other binding partner. The interaction is dependent upon the presence of a particular structure of the protein, e.g., the antigenic determinant or epitope, recognized by the binding molecule. For example, if an antibody is specific for epitope "A," the presence of a polypeptide containing epitope A, or the presence of free unlabeled A, in a reaction containing free labeled A and the antibody will reduce the amount of labeled A that binds to the antibody.

"Substitution" refers to the replacement of at least one nucleotide or amino acid by a different nucleotide or amino acid.

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"Substrate" refers to any suitable rigid or semi-rigid support including, e.g., membranes, filters, chips, slides, wafers, fibers, magnetic or nonmagnetic beads, gels, tubing, plates, polymers, microparticles or capillaries. The substrate can have a variety of surface forms, such as wells, trenches, pins, channels and pores, to which polynucleotides or polypeptides are bound.

A "transcript image" refers to the collective pattern of gene expression by a particular tissue or cell type under given conditions at a given time.

"Transformation" refers to a process by which exogenous DNA enters a recipient cell.

Transformation may occur under natural or artificial conditions using various methods well known in the art. Transformation may rely on any known method for the insertion of foreign nucleic acid sequences into a prokaryotic or eukaryotic host cell. The method is selected based on the host cell being transformed.

"Transformants" include stably transformed cells in which the inserted DNA is capable of replication either as an autonomously replicating plasmid or as part of the host chromosome, as well as cells which transiently express inserted DNA or RNA.

A "transgenic organism," as used herein, is any organism, including but not limited to animals and plants, in which one or more of the cells of the organism contains heterologous nucleic acid introduced by way of human intervention, such as by transgenic techniques well known in the art. The nucleic acid is introduced into the cell, directly or indirectly by introduction into a precursor of the cell, by way of deliberate genetic manipulation, such as by microinjection or by infection with a recombinant virus. The term genetic manipulation does not include classical cross-breeding, or in vitro fertilization, but rather is directed to the introduction of a recombinant DNA molecule. The transgenic organisms contemplated in accordance with the present invention include bacteria, cyanobacteria, fungi, and plants and animals. The isolated DNA of the present invention can be introduced into the host by methods known in the art, for example infection, transfection, transformation or transconjugation. Techniques for transferring the DNA of the present invention into such organisms are widely known and provided in references such as Sambrook et al. (1989), supra.

VA). Prior to mRNA isolation, cell lines were untreated, treated with a pharmaceutical agent such as 5'-aza-2'-deoxycytidine, treated with an activating agent such as lipopolysaccharide in the case of leukocytic cell lines, or, in the case of endothelial cell lines, subjected to shear stress.

Sequencing of the cDNAs

Methods for DNA sequencing are well known in the art. Conventional enzymatic methods employ the Klenow fragment of DNA polymerase I, SEQUENASE DNA polymerase (U.S. Biochemical Corporation, Cleveland OH), Taq polymerase (PE Biosystems, Foster City CA), thermostable T7 polymerase (Amersham Pharmacia Biotech, Inc. (Amersham Pharmacia Biotech), Piscataway NJ), or combinations of polymerases and proofreading exonucleases such as those found in the ELONGASE amplification system (Life Technologies Inc. (Life Technologies), Gaithersburg MD), to extend the nucleic acid sequence from an oligonucleotide primer annealed to the DNA template of interest. Methods have been developed for the use of both single-stranded and double-stranded templates. Chain termination reaction products may be electrophoresed on urea-polyacrylamide gels and detected either by autoradiography (for radioisotope-labeled nucleotides) or by fluorescence (for fluorophore-labeled nucleotides). Automated methods for mechanized reaction preparation, sequencing, and analysis using fluorescence detection methods have been developed. Machines used to prepare cDNAs for sequencing can include the MICROLAB 2200 liquid transfer system (Hamilton Company (Hamilton), Reno NV), Peltier thermal cycler (PTC200; MJ Research, Inc. (MJ Research), Watertown MA), and ABI CATALYST 800 thermal cycler (PE Biosystems). Sequencing can be carried out using, for example, the ABI 373 or 377 (PE Biosystems) or MEGABACE 1000 (Molecular Dynamics, Inc. (Molecular Dynamics), Sunnyvale CA) DNA sequencing systems, or other automated and manual sequencing systems well known in the art.

The nucleotide sequences of the Sequence Listing have been prepared by current, state-of-the-art, automated methods and, as such, may contain occasional sequencing errors or unidentified nucleotides. Such unidentified nucleotides are designated by an N. These infrequent unidentified bases do not represent a hindrance to practicing the invention for those skilled in the art. Several methods employing standard recombinant techniques may be used to correct errors and complete the missing sequence information. (See, e.g., those described in Ausubel, F.M. et al. (1997) Short Protocols in Molecular Biology, John Wiley & Sons, New York NY; and Sambrook, J. et al. (1989) Molecular Cloning, A Laboratory Manual, Cold Spring Harbor Press, Plainview NY.)

Assembly of cDNA Sequences

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Human polynucleotide sequences may be assembled using programs or algorithms well known in the art. Sequences to be assembled are related, wholly or in part, and may be derived from a single or many different transcripts. Assembly of the sequences can be performed using such programs as

A "variant" of a particular polypeptide sequence is defined as a polypeptide sequence having at least 40% sequence identity to the particular polypeptide sequence over a certain length of one of the polypeptide sequences using blastp with the "BLAST 2 Sequences" tool Version 2.0.9 (May-07-1999) set at default parameters. Such a pair of polypeptides may show, for example, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, at least 95%, or at least 98% or greater sequence identity over a certain defined length of one of the polypeptides. THE INVENTION

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In a particular embodiment, cDNA sequences derived from human tissues and cell lines were aligned based on nucleotide sequence identity and assembled into "consensus" or "template" sequences which are designated by the template identification numbers (template IDs) in column 2 of Table 1. The sequence identification numbers (SEQ ID NO:s) corresponding to the template IDs are shown in column 1. Segments of the template sequences are defined by the "start" and "stop" nucleotide positions listed in columns 3 and 4. These segments, when translated in the reading frames indicated in column 5, have similarity to signal peptide (SP) or transmembrane (TM) domain consensus sequences, as indicated in column 6.

The invention incorporates the nucleic acid sequences of these templates as disclosed in the Sequence Listing and the use of these sequences in the diagnosis and treatment of disease states characterized by defects in cell signaling. The invention further utilizes these sequences in hybridization and amplification technologies, and in particular, in technologies which assess gene expression patterns correlated with specific cells or tissues and their responses in vivo or in vitro to pharmaceutical agents, toxins, and other treatments. In this manner, the sequences of the present invention are used to develop a transcript image for a particular cell or tissue.

Derivation of Nucleic Acid Sequences

cDNA was isolated from libraries constructed using RNA derived from normal and diseased human tissues and cell lines. The human tissues and cell lines used for cDNA library construction were selected from a broad range of sources to provide a diverse population of cDNAs representative of gene transcription throughout the human body. Descriptions of the human tissues and cell lines used for cDNA library construction are provided in the LIFESEQ database (Incyte Genomics, Inc. (Incyte), Palo Alto CA). Human tissues were broadly selected from, for example, cardiovascular, dermatologic, endocrine, gastrointestinal, hematopoietic/immune system, musculoskeletal, neural, reproductive, and 30

Cell lines used for cDNA library construction were derived from, for example, leukemic cells, teratocarcinomas, neuroepitheliomas, cervical carcinoma, lung fibroblasts, and endothelial cells. Such cell lines include, for example, THP-1, Jurkat, HUVEC, hNT2, WI38, HeLa, and other cell lines commonly used and available from public depositories (American Type Culture Collection, Manassas

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Computer programs known to those of skill in the art for performing computer-assisted searches for amino acid and nucleic acid sequence similarity, include, for example, Basic Local Alignment Search Tool (BLAST; Altschul, S.F. (1993) J. Mol. Evol. 36:290-300; Altschul, S.F. et al. (1990) J. Mol. Biol. 215:403-410). BLAST is especially useful in determining exact matches and comparing two sequence fragments of arbitrary but equal lengths, whose alignment is locally maximal and for which the alignment score meets or exceeds a threshold or cutoff score set by the user (Karlin, S. et al. (1988) Proc. Natl. Acad. Sci. USA 85:841-845). Using an appropriate search tool (e.g., BLAST or HMM), GenBank, SwissProt, BLOCKS, PFAM and other databases may be searched for sequences containing regions of homology to a query sptm or SPTM of the present invention.

Other approaches to the identification, assembly, storage, and display of nucleotide and polypeptide sequences are provided in "Relational Database for Storing Biomolecule Information," U.S.S.N. 08/947,845, filed October 9, 1997; "Project-Based Full-Length Biomolecular Sequence Database," U.S.S.N. 08/811,758, filed March 6, 1997; and "Relational Database and System for Storing Information Relating to Biomolecular Sequences," U.S.S.N. 09/034,807, filed March 4, 1998, all of which are incorporated by reference herein in their entirety.

Protein hierarchies can be assigned to the putative encoded polypeptide based on, e.g., motif, BLAST, or biological analysis. Methods for assigning these hierarchies are described, for example, in "Database System Employing Protein Function Hierarchies for Viewing Biomolecular Sequence Data," U.S.S.N. 08/812,290, filed March 6, 1997, incorporated herein by reference.

20 Human Secretory Sequences

The sptm of the present invention may be used for a variety of diagnostic and therapeutic purposes. For example, an sptm may be used to diagnose a particular condition, disease, or disorder associated with cell signaling. Such conditions, diseases, and disorders include, but are not limited to, a cell proliferative disorder such as actinic keratosis, arteriosclerosis, atherosclerosis, bursitis, cirrhosis, hepatitis, mixed connective tissue disease (MCTD), myelofibrosis, paroxysmal nocturnal hemoglobinuria, polycythemia vera, psoriasis, primary thrombocythemia, and cancers including adenocarcinoma, leukemia, lymphoma, melanoma, myeloma, sarcoma, teratocarcinoma, and, in particular, a cancer of the adrenal gland, bladder, bone, bone marrow, brain, breast, cervix, gall bladder, ganglia, gastrointestinal tract, heart, kidney, liver, lung, muscle, ovary, pancreas, parathyroid, penis, prostate, salivary glands, skin, spleen, testis, thymus, thyroid, and uterus; an immune system disorder such as such as inflammation, actinic keratosis, acquired immunodeficiency syndrome (AIDS), Addison's disease, adult respiratory distress syndrome, allergies, ankylosing spondylitis, amyloidosis, anemia, arteriosclerosis, asthma, atherosclerosis, autoimmune hemolytic anemia, autoimmune thyroiditis, bronchitis, bursitis, cholecystitis, cirrhosis, contact dermatitis, Crohn's disease, atopic dermatitis, dermatomyositis, diabetes mellitus, emphysema, erythroblastosis fetalis,

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PHRAP (Phils Revised Assembly Program) and the GELVIEW fragment assembly system (GCG), or other methods known in the art.

Alternatively, cDNA sequences are used as "component" sequences that are assembled into "template" or "consensus" sequences as follows. Sequence chromatograms are processed, verified, and quality scores are obtained using PHRED. Raw sequences are edited using an editing pathway known as Block 1 (See, e.g., the LIFESEQ Assembled User Guide, Incyte Genomics, Palo Alto, CA). A series of BLAST comparisons is performed and low-information segments and repetitive elements (e.g., dinucleotide repeats, Alu repeats, etc.) are replaced by "n's", or masked, to prevent spurious matches. Mitochondrial and ribosomal RNA sequences are also removed. The processed sequences are then loaded into a relational database management system (RDMS) which assigns edited sequences to existing templates, if available. When additional sequences are added into the RDMS, a process is initiated which modifies existing templates or creates new templates from works in progress (i.e., nonfinal assembled sequences) containing queued sequences or the sequences themselves. After the new sequences have been assigned to templates, the templates can be merged into bins. If multiple templates exist in one bin, the bin can be split and the templates reannotated.

Once gene bins have been generated based upon sequence alignments, bins are "clone joined" based upon clone information. Clone joining occurs when the 5' sequence of one clone is present in one bin and the 3' sequence from the same clone is present in a different bin, indicating that the two bins should be merged into a single bin. Only bins which share at least two different clones are merged.

A resultant template sequence may contain either a partial or a full length open reading frame, 20 or all or part of a genetic regulatory element. This variation is due in part to the fact that the full length cDNAs of many genes are several hundred, and sometimes several thousand, bases in length. With current technology, cDNAs comprising the coding regions of large genes cannot be cloned because of vector limitations, incomplete reverse transcription of the mRNA, or incomplete "second strand" synthesis. Template sequences may be extended to include additional contiguous sequences derived from the parent RNA transcript using a variety of methods known to those of skill in the art. Extension may thus be used to achieve the full length coding sequence of a gene. Analysis of the cDNA Sequences

The cDNA sequences are analyzed using a variety of programs and algorithms which are well known in the art. (See, e.g., Ausubel, 1997, supra, Chapter 7.7; Meyers, R.A. (Ed.) (1995) Molecular 30 Biology and Biotechnology, Wiley VCH, New York NY, pp. 856-853; and Table 4.) These analyses comprise both reading frame determinations, e.g., based on triplet codon periodicity for particular organisms (Fickett, J.W. (1982) Nucleic Acids Res. 10:5303-5318); analyses of potential start and stop codons; and homology searches.

of analysis is useful, for example, to assess the relative levels of sptm expression in fully or partially differentiated cells or tissues, to determine if changes in sptm expression levels are correlated with the development or progression of specific disease states, and to assess the response of a cell or tissue to a specific therapy, for example, in pharmacological or toxicological studies. Methods for the analysis of sptm expression are based on hybridization and amplification technologies and include membrane-based procedures such as northern blot analysis, high-throughput procedures that utilize, for example, microarrays, and PCR-based procedures.

Hybridization and Genetic Analysis

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The sptm, their fragments, or complementary sequences, may be used to identify the presence of and/or to determine the degree of similarity between two (or more) nucleic acid sequences. The sptm may be hybridized to naturally occurring or recombinant nucleic acid sequences under appropriately selected temperatures and salt concentrations. Hybridization with a probe based on the nucleic acid sequence of at least one of the sptm allows for the detection of nucleic acid sequences, including genomic sequences, which are identical or related to the sptm of the Sequence Listing. Probes may be selected from non-conserved or unique regions of at least one of the polynucleotides of SEQ ID NO:1-63 and tested for their ability to identify or amplify the target nucleic acid sequence using standard protocols.

Polynucleotide sequences that are capable of hybridizing, in particular, to those shown in SEQ ID NO:1-63 and fragments thereof, can be identified using various conditions of stringency. (See, e.g., Wahl, G.M. and S.L. Berger (1987) Methods Enzymol. 152:399-407; Kimmel, A.R. (1987) Methods Enzymol. 152:507-511.) Hybridization conditions are discussed in "Definitions."

A probe for use in Southern or northern hybridization may be derived from a fragment of an sptm sequence, or its complement, that is up to several hundred nucleotides in length and is either single-stranded or double-stranded. Such probes may be hybridized in solution to biological materials such as plasmids, bacterial, yeast, or human artificial chromosomes, cleared or sectioned tissues, or to artificial substrates containing sptm. Microarrays are particularly suitable for identifying the presence of and detecting the level of expression for multiple genes of interest by examining gene expression correlated with, e.g., various stages of development, treatment with a drug or compound, or disease progression. An array analogous to a dot or slot blot may be used to arrange and link polynucleotides to the surface of a substrate using one or more of the following: mechanical (vacuum), chemical, thermal, or UV bonding procedures. Such an array may contain any number of sptm and may be produced by hand or by using available devices, materials, and machines.

Microarrays may be prepared, used, and analyzed using methods known in the art. (See, e.g., Brennan, T.M. et al. (1995) U.S. Patent No. 5,474,796; Schena, M. et al. (1996) Proc. Natl. Acad. Sci.

erythema nodosum, atrophic gastritis, glomerulonephritis, Goodpasture's syndrome, gout, Graves' disease, Hashimoto's thyroiditis, paroxysmal nocturnal hemoglobinuria, hepatitis, hypereosinophilia, irritable bowel syndrome, episodic lymphopenia with lymphocytotoxins, mixed connective tissue disease (MCTD), multiple sclerosis, myasthenia gravis, myocardial or pericardial inflammation, myelofibrosis, osteoarthritis, osteoporosis, pancreatitis, polycythemia vera, polymyositis, psoriasis, Reiter's syndrome, rheumatoid arthritis, scleroderma, Sjögren's syndrome, systemic anaphylaxis, systemic lupus erythematosus, systemic sclerosis, primary thrombocythemia, thrombocytopenic purpura, ulcerative colitis, uveitis, Werner syndrome, complications of cancer, hemodialysis, and extracorporeal circulation, trauma, and hematopoietic cancer including lymphoma, leukemia, and myeloma; and a neurological disorder such as epilepsy, ischemic cerebrovascular disease, stroke, 10 cerebral neoplasms, Alzheimer's disease, Pick's disease, Huntington's disease, dementia, Parkinson's disease and other extrapyramidal disorders, amyotrophic lateral sclerosis and other motor neuron disorders, progressive neural muscular atrophy, retinitis pigmentosa, hereditary ataxias, multiple sclerosis and other demyelinating diseases, bacterial and viral meningitis, brain abscess, subdural empyema, epidural abscess, suppurative intracranial thrombophlebitis, myelitis and radiculitis, viral 15 central nervous system disease, prion diseases including kuru, Creutzfeldt-Jakob disease, and Gerstmann-Straussler-Scheinker syndrome, fatal familial insomnia, nutritional and metabolic diseases of the nervous system, neurofibromatosis, tuberous sclerosis, cerebelloretinal hemangioblastomatosis, encephalotrigeminal syndrome, mental retardation and other developmental disorder of the central nervous system, cerebral palsy, a neuroskeletal disorder, an autonomic nervous system disorder, a cranial nerve disorder, a spinal cord disease, muscular dystrophy and other neuromuscular disorder, a peripheral nervous system disorder, dermatomyositis and polymyositis, inherited, metabolic, endocrine, and toxic myopathy, myasthenia gravis, periodic paralysis, a mental disorder including mood, anxiety, and schizophrenic disorder, seasonal affective disorder (SAD), akathesia, amnesia, catatonia, diabetic neuropathy, tardive dyskinesia, dystonias, paranoid psychoses, postherpetic neuralgia, and Tourette's disorder. The sptm can be used to detect the presence of, or to quantify the amount of, an sptm-related polynucleotide in a sample. This information is then compared to information obtained from appropriate reference samples, and a diagnosis is established. Alternatively, a polynucleotide complementary to a given sptm can inhibit or inactivate a therapeutically relevant gene related to the sptm.

Analysis of sptm Expression Patterns 30

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The expression of sptm may be routinely assessed by hybridization-based methods to determine, for example, the tissue-specificity, disease-specificity, or developmental stage-specificity of sptm expression. For example, the level of expression of sptm may be compared among different cell types or tissues, among diseased and normal cell types or tissues, among cell types or tissues at different developmental stages, or among cell types or tissues undergoing various treatments. This type

linkage maps can be found in various scientific journals or at the Online Mendelian Inheritance in Man (OMIM) World Wide Web site.

In another embodiment of the invention, sptm sequences may be used to generate hybridization probes useful in chromosomal mapping of naturally occurring genomic sequences. Either coding or noncoding sequences of sptm may be used, and in some instances, noncoding sequences may be preferable over coding sequences. For example, conservation of an sptm coding sequence among members of a multi-gene family may potentially cause undesired cross hybridization during chromosomal mapping. The sequences may be mapped to a particular chromosome, to a specific region of a chromosome, or to artificial chromosome constructions, e.g., human artificial chromosomes (HACs), yeast artificial chromosomes (YACs), bacterial artificial chromosomes (BACs), bacterial P1 constructions, or single chromosome cDNA libraries. (See, e.g., Harrington, J.J. et al. (1997) Nat. Genet. 15:345-355; Price, C.M. (1993) Blood Rev. 7:127-134; and Trask, B.J. (1991) Trends Genet. 7:149-154.)

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Fluorescent <u>in situ</u> hybridization (FISH) may be correlated with other physical chromosome mapping techniques and genetic map data. (See, e.g., Meyers, <u>supra</u>, pp. 965-968.) Correlation between the location of sptm on a physical chromosomal map and a specific disorder, or a predisposition to a specific disorder, may help define the region of DNA associated with that disorder. The sptm sequences may also be used to detect polymorphisms that are genetically linked to the inheritance of a particular condition, disease, or disorder.

In situ hybridization of chromosomal preparations and genetic mapping techniques, such as linkage analysis using established chromosomal markers, may be used for extending existing genetic maps. Often the placement of a gene on the chromosome of another mammalian species, such as mouse, may reveal associated markers even if the number or arm of the corresponding human chromosome is not known. These new marker sequences can be mapped to human chromosomes and may provide valuable information to investigators searching for disease genes using positional cloning or other gene discovery techniques. Once a disease or syndrome has been crudely correlated by genetic linkage with a particular genomic region, e.g., ataxia-telangiectasia to 11q22-23, any sequences mapping to that area may represent associated or regulatory genes for further investigation. (See, e.g., Gatti, R.A. et al. (1988) Nature 336:577-580.) The nucleotide sequences of the subject invention may also be used to detect differences in chromosomal architecture due to translocation, inversion, etc., among normal, carrier, or affected individuals.

Once a disease-associated gene is mapped to a chromosomal region, the gene must be cloned in order to identify mutations or other alterations (e.g., translocations or inversions) that may be correlated with disease. This process requires a physical map of the chromosomal region containing the disease-gene of interest along with associated markers. A physical map is necessary for determining the

USA 93:10614-10619; Baldeschweiler et al. (1995) PCT application WO95/251116; Shalon, D. et al. (1995) PCT application WO95/35505; Heller, R.A. et al. (1997) Proc. Natl. Acad. Sci. USA 94:2150-2155; and Heller, M.J. et al. (1997) U.S. Patent No. 5,605,662.)

Probes may be labeled by either PCR or enzymatic techniques using a variety of commercially available reporter molecules. For example, commercial kits are available for radioactive and chemiluminescent labeling (Amersham Pharmacia Biotech) and for alkaline phosphatase labeling (Life Technologies). Alternatively, sptm may be cloned into commercially available vectors for the production of RNA probes. Such probes may be transcribed in the presence of at least one labeled nucleotide (e.g., ³²P-ATP, Amersham Pharmacia Biotech).

Additionally the polynucleotides of SEQ ID NO:1-63 or suitable fragments thereof can be used to isolate full length cDNA sequences utilizing hybridization and/or amplification procedures well known in the art, e.g., cDNA library screening, PCR amplification, etc. The molecular cloning of such full length cDNA sequences may employ the method of cDNA library screening with probes using the hybridization, stringency, washing, and probing strategies described above and in Ausubel, <u>supra</u>,

15 Chapters 3, 5, and 6. These procedures may also be employed with genomic libraries to isolate genomic sequences of sptm in order to analyze, e.g., regulatory elements.

Genetic Mapping

Gene identification and mapping are important in the investigation and treatment of almost all conditions, diseases, and disorders. Cancer, cardiovascular disease, Alzheimer's disease, arthritis, diabetes, and mental illnesses are of particular interest. Each of these conditions is more complex than the single gene defects of sickle cell anemia or cystic fibrosis, with select groups of genes being predictive of predisposition for a particular condition, disease, or disorder. For example, cardiovascular disease may result from malfunctioning receptor molecules that fail to clear cholesterol from the bloodstream, and diabetes may result when a particular individual's immune system is activated by an infection and attacks the insulin-producing cells of the pancreas. In some studies, Alzheimer's disease has been linked to a gene on chromosome 21; other studies predict a different gene and location. Mapping of disease genes is a complex and reiterative process and generally proceeds from genetic linkage analysis to physical mapping.

As a condition is noted among members of a family, a genetic linkage map traces parts of chromosomes that are inherited in the same pattern as the condition. Statistics link the inheritance of particular conditions to particular regions of chromosomes, as defined by RFLP or other markers. (See, for example, Lander, E. S. and Botstein, D. (1986) Proc. Natl. Acad. Sci. USA 83:7353-7357.) Occasionally, genetic markers and their locations are known from previous studies. More often, however, the markers are simply stretches of DNA that differ among individuals. Examples of genetic

primers for amplifying and isolating such selected DNA, which can then be sequenced. Using this technique, an individual can be identified through a unique set of DNA sequences. Once a unique ID database is established for an individual, positive identification of that individual can be made from extremely small tissue samples.

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In a particular aspect, oligonucleotide primers derived from the sptm of the invention may be used to detect single nucleotide polymorphisms (SNPs). SNPs are substitutions, insertions and deletions that are a frequent cause of inherited or acquired genetic disease in humans. Methods of SNP detection include, but are not limited to, single-stranded conformation polymorphism (SSCP) and fluorescent SSCP (fSSCP) methods. In SSCP, oligonucleotide primers derived from sptm are used to amplify DNA using the polymerase chain reaction (PCR). The DNA may be derived, for example, from diseased or normal tissue, biopsy samples, bodily fluids, and the like. SNPs in the DNA cause differences in the secondary and tertiary structures of PCR products in single-stranded form, and these differences are detectable using gel electrophoresis in non-denaturing gels. In fSCCP, the oligonucleotide primers are fluorescently labeled, which allows detection of the amplimers in highthroughput equipment such as DNA sequencing machines. Additionally, sequence database analysis methods, termed in silico SNP (isSNP), are capable of identifying polymorphisms by comparing the sequences of individual overlapping DNA fragments which assemble into a common consensus sequence. These computer-based methods filter out sequence variations due to laboratory preparation of DNA and sequencing errors using statistical models and automated analyses of DNA sequence chromatograms. In the alternative, SNPs may be detected and characterized by mass spectrometry using, for example, the high throughput MASSARRAY system (Sequenom, Inc., San Diego CA).

DNA-based identification techniques are critical in forensic technology. DNA sequences taken from very small biological samples such as tissues, e.g., hair or skin, or body fluids, e.g., blood, saliva, semen, etc., can be amplified using, e.g., PCR, to identify individuals. (See, e.g., Erlich, H. (1992) PCR Technology, Freeman and Co., New York, NY). Similarly, polynucleotides of the present invention can be used as polymorphic markers.

There is also a need for reagents capable of identifying the source of a particular tissue. Appropriate reagents can comprise, for example, DNA probes or primers prepared from the sequences of the present invention that are specific for particular tissues. Panels of such reagents can identify tissue by species and/or by organ type. In a similar fashion, these reagents can be used to screen tissue cultures for contamination.

The polynucleotides of the present invention can also be used as molecular weight markers on nucleic acid gels or Southern blots, as diagnostic probes for the presence of a specific mRNA in a particular cell type, in the creation of subtracted cDNA libraries which aid in the discovery of novel

nucleotide sequence of and order of marker genes on a particular chromosomal region. Physical mapping techniques are well known in the art and require the generation of overlapping sets of cloned DNA fragments from a particular organelle, chromosome, or genome. These clones are analyzed to reconstruct and catalog their order. Once the position of a marker is determined, the DNA from that region is obtained by consulting the catalog and selecting clones from that region. The gene of interest is located through positional cloning techniques using hybridization or similar methods. Diagnostic Uses

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The sptm of the present invention may be used to design probes useful in diagnostic assays. Such assays, well known to those skilled in the art, may be used to detect or confirm conditions, disorders, or diseases associated with abnormal levels of sptm expression. Labeled probes developed 10 from sptm sequences are added to a sample under hybridizing conditions of desired stringency. In some instances, sptm, or fragments or oligonucleotides derived from sptm, may be used as primers in amplification steps prior to hybridization. The amount of hybridization complex formed is quantified and compared with standards for that cell or tissue. If sptm expression varies significantly from the standard, the assay indicates the presence of the condition, disorder, or disease. Qualitative or quantitative diagnostic methods may include northern, dot blot, or other membrane or dip-stick based technologies or multiple-sample format technologies such as PCR, enzyme-linked immunosorbent assay (ELISA)-like, pin, or chip-based assays.

The probes described above may also be used to monitor the progress of conditions, disorders, or diseases associated with abnormal levels of sptm expression, or to evaluate the efficacy of a 20 particular therapeutic treatment. The candidate probe may be identified from the sptm that are specific to a given human tissue and have not been observed in GenBank or other genome databases. Such a probe may be used in animal studies, preclinical tests, clinical trials, or in monitoring the treatment of an individual patient. In a typical process, standard expression is established by methods well known in the art for use as a basis of comparison, samples from patients affected by the disorder or disease are combined with the probe to evaluate any deviation from the standard profile, and a therapeutic agent is administered and effects are monitored to generate a treatment profile. Efficacy is evaluated by determining whether the expression progresses toward or returns to the standard normal pattern. Treatment profiles may be generated over a period of several days or several months. Statistical methods well known to those skilled in the art may be use to determine the significance of such 30

The polynucleotides are also useful for identifying individuals from minute biological samples, for example, by matching the RFLP pattern of a sample's DNA to that of an individual's DNA. The polynucleotides of the present invention can also be used to determine the actual base-by-base DNA sequence of selected portions of an individual's genome. These sequences can be used to prepare PCR

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bound molecule. Examples of such molecules include antibodies, oligonucleotides, proteins (e.g., receptors), or small molecules.

Preferably, the molecule is closely related to the natural ligand of the polypeptide, e.g., a ligand or fragment thereof, a natural substrate, or a structural or functional mimetic. (See, Coligan et al., (1991) Current Protocols in Immunology 1(2): Chapter 5.) Similarly, the molecule can be closely related to the natural receptor to which the polypeptide binds, or to at least a fragment of the receptor, e.g., the active site. In either case, the molecule can be rationally designed using known techniques. Preferably, the screening for these molecules involves producing appropriate cells which express the polypeptide, either as a secreted protein or on the cell membrane. Preferred cells include cells from mammals, yeast, Drosophila, or E. coli. Cells expressing the polypeptide or cell membrane fractions which contain the expressed polypeptide are then contacted with a test compound and binding, stimulation, or inhibition of activity of either the polypeptide or the molecule is analyzed.

An assay may simply test binding of a candidate compound to the polypeptide, wherein binding is detected by a fluorophore, radioisotope, enzyme conjugate, or other detectable label. Alternatively, the assay may assess binding in the presence of a labeled competitor.

Additionally, the assay can be carried out using cell-free preparations, polypeptide/molecule affixed to a solid support, chemical libraries, or natural product mixtures. The assay may also simply comprise the steps of mixing a candidate compound with a solution containing a polypeptide, measuring polypeptide/molecule activity or binding, and comparing the polypeptide/molecule activity or binding to a standard.

Preferably, an ELISA assay using, e.g., a monoclonal or polyclonal antibody, can measure polypeptide level in a sample. The antibody can measure polypeptide level by either binding, directly or indirectly, to the polypeptide or by competing with the polypeptide for a substrate.

All of the above assays can be used in a diagnostic or prognostic context. The molecules discovered using these assays can be used to treat disease or to bring about a particular result in a patient (e.g., blood vessel growth) by activating or inhibiting the polypeptide/molecule. Moreover, the assays can discover agents which may inhibit or enhance the production of the polypeptide from suitably manipulated cells or tissues.

Transcript Imaging and Toxicological Testing

Another embodiment relates to the use of sptm to develop a transcript image of a tissue or cell type. A transcript image represents the global pattern of gene expression by a particular tissue or cell type. Global gene expression patterns are analyzed by quantifying the number of expressed genes and their relative abundance under given conditions and at a given time. (See Seilhamer et al., "Comparative Gene Transcript Analysis," U.S. Patent Number 5,840,484, expressly incorporated by reference herein.) Thus a transcript image may be generated by hybridizing the polynucleotides of the

polynucleotides, in selection and synthesis of oligomers for attachment to an array or other support, and as an antigen to elicit an immune response. Disease Model Systems Using SPTM

The polynucleotides encoding SPTM or their mammalian homologs may be "knocked out" in an animal model system using homologous recombination in embryonic stem (ES) cells. Such techniques are well known in the art and are useful for the generation of animal models of human disease. (See, e.g., U.S. Patent Number 5,175,383 and U.S. Patent Number 5,767,337.) For example, mouse ES cells, such as the mouse 129/SvJ cell line, are derived from the early mouse embryo and grown in culture. The ES cells are transformed with a vector containing the gene of interest disrupted by a marker gene, e.g., the neomycin phosphotransferase gene (neo; Capecchi, M.R. (1989) Science 244:1288-1292). The vector integrates into the corresponding region of the host genome by homologous recombination. Alternatively, homologous recombination takes place using the Cre-loxP system to knockout a gene of interest in a tissue- or developmental stage-specific manner (Marth, J.D. (1996) Clin. Invest. 97:1999-2002; Wagner, K.U. et al. (1997) Nucleic Acids Res. 25:4323-4330).

Transformed ES cells are identified and microinjected into mouse cell blastocysts such as those from the C57BL/6 mouse strain. The blastocysts are surgically transferred to pseudopregnant dams, and the resulting chimeric progeny are genotyped and bred to produce heterozygous or homozygous strains. Transgenic animals thus generated may be tested with potential therapeutic or toxic agents.

Polynucleotides encoding SPTM may also be manipulated in vitro in ES cells derived from human blastocysts. Human ES cells have the potential to differentiate into at least eight separate cell 20 lineages including endoderm, mesoderm, and ectodermal cell types. These cell lineages differentiate into, for example, neural cells, hematopoietic lineages, and cardiomyocytes (Thomson, J.A. et al. (1998)

Polynucleotides encoding SPTM can also be used to create "knockin" humanized animals (pigs) or transgenic animals (mice or rats) to model human disease. With knockin technology, a region 25 of sptm is injected into animal ES cells, and the injected sequence integrates into the animal cell genome. Transformed cells are injected into blastulae, and the blastulae are implanted as described above. Transgenic progeny or inbred lines are studied and treated with potential pharmaceutical agents to obtain information on treatment of a human disease. Alternatively, a mammal inbred to overexpress sptm, resulting, e.g., in the secretion of SPTM in its milk, may also serve as a convenient source of that 30 protein (Janne, J. et al. (1998) Biotechnol. Annu. Rev. 4:55-74). Screening Assays

SPTM encoded by polynucleotides of the present invention may be used to screen for molecules that bind to or are bound by the encoded polypeptides. The binding of the polypeptide and the molecule may activate (agonist), increase, inhibit (antagonist), or decrease activity of the polypeptide or the 35

Another particular embodiment relates to the use of SPTM encoded by polynucleotides of the present invention to analyze the proteome of a tissue or cell type. The term proteome refers to the global pattern of protein expression in a particular tissue or cell type. Each protein component of a proteome can be subjected individually to further analysis. Proteome expression patterns, or profiles, are analyzed by quantifying the number of expressed proteins and their relative abundance under given conditions and at a given time. A profile of a cell's proteome may thus be generated by separating and analyzing the polypeptides of a particular tissue or cell type. In one embodiment, the separation is achieved using two-dimensional gel electrophoresis, in which proteins from a sample are separated by isoelectric focusing in the first dimension, and then according to molecular weight by sodium dodecyl sulfate slab gel electrophoresis in the second dimension (Steiner and Anderson, supra). The proteins are visualized in the gel as discrete and uniquely positioned spots, typically by staining the gel with an agent such as Coomassie Blue or silver or fluorescent stains. The optical density of each protein spot is generally proportional to the level of the protein in the sample. The optical densities of equivalently positioned protein spots from different samples, for example, from biological samples either treated or untreated with a test compound or therapeutic agent, are compared to identify any changes in protein spot density related to the treatment. The proteins in the spots are partially sequenced using, for example, standard methods employing chemical or enzymatic cleavage followed by mass spectrometry. The identity of the protein in a spot may be determined by comparing its partial sequence, preferably of at least 5 contiguous amino acid residues, to the polypeptide sequences of the present invention. In some cases, further sequence data may be obtained for definitive protein identification.

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A proteomic profile may also be generated using antibodies specific for SPTM to quantify the levels of SPTM expression. In one embodiment, the antibodies are used as elements on a microarray, and protein expression levels are quantified by exposing the microarray to the sample and detecting the levels of protein bound to each array element (Lucking, A. et al. (1999) Anal. Biochem. 270:103-11; Mendoze, L. G. et al. (1999) Biotechniques 27:778-88). Detection may be performed by a variety of methods known in the art, for example, by reacting the proteins in the sample with a thiol- or amino-reactive fluorescent compound and detecting the amount of fluorescence bound at each array element.

Toxicant signatures at the proteome level are also useful for toxicological screening, and should be analyzed in parallel with toxicant signatures at the transcript level. There is a poor correlation between transcript and protein abundances for some proteins in some tissues (Anderson, N. L. and Seilhamer, J. (1997) Electrophoresis 18:533-537), so proteome toxicant signatures may be useful in the analysis of compounds which do not significantly affect the transcript image, but which alter the proteomic profile. In addition, the analysis of transcripts in body fluids is difficult, due to rapid degradation of mRNA, so proteomic profiling may be more reliable and informative in such cases.

present invention or their complements to the totality of transcripts or reverse transcripts of a particular tissue or cell type. In one embodiment, the hybridization takes place in high-throughput format, wherein the polynucleotides of the present invention or their complements comprise a subset of a plurality of elements on a microarray. The resultant transcript image would provide a profile of gene activity pertaining to cell signaling.

Transcript images which profile sptm expression may be generated using transcripts isolated from tissues, cell lines, biopsies, or other biological samples. The transcript image may thus reflect sptm expression in vivo, as in the case of a tissue or biopsy sample, or in vitro, as in the case of a cell line.

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10 Transcript images which profile sptm expression may also be used in conjunction with in vitro model systems and preclinical evaluation of pharmaceuticals, as well as toxicological testing of industrial and naturally-occurring environmental compounds. All compounds induce characteristic gene expression patterns, frequently termed molecular fingerprints or toxicant signatures, which are indicative of mechanisms of action and toxicity (Nuwaysir, E. F. et al. (1999) Mol. Carcinog. 24:153-159; Steiner, S. and Anderson, N. L. (2000) Toxicol. Lett. 112-113:467-71, expressly incorporated by 15 reference herein). If a test compound has a signature similar to that of a compound with known toxicity, it is likely to share those toxic properties. These fingerprints or signatures are most useful and refined when they contain expression information from a large number of genes and gene families. Ideally, a genome-wide measurement of expression provides the highest quality signature. Even genes whose expression is not altered by any tested compounds are important as well, as the levels of 20 expression of these genes are used to normalize the rest of the expression data. The normalization procedure is useful for comparison of expression data after treatment with different compounds. While the assignment of gene function to elements of a toxicant signature aids in interpretation of toxicity mechanisms, knowledge of gene function is not necessary for the statistical matching of signatures which leads to prediction of toxicity. (See, for example, Press Release 00-02 from the National 25 Institute of Environmental Health Sciences, released February 29, 2000, available at http://www.niehs.nih.gov/oc/news/toxchip.htm.) Therefore, it is important and desirable in toxicological screening using toxicant signatures to include all expressed gene sequences.

In one embodiment, the toxicity of a test compound is assessed by treating a biological sample containing nucleic acids with the test compound. Nucleic acids that are expressed in the treated biological sample are hybridized with one or more probes specific to the polynucleotides of the present invention, so that transcript levels corresponding to the polynucleotides of the present invention may be quantified. The transcript levels in the treated biological sample are compared with levels in an untreated biological sample. Differences in the transcript levels between the two samples are indicative of a toxic response caused by the test compound in the treated sample.

al. (1997) Biochem. Mol. Med. 62(1):11-22.) An antisense sequence is a polynucleotide sequence capable of specifically hybridizing to at least a portion of the target sequence. Antisense sequences bind to cellular mRNA and/or genomic DNA, affecting translation and/or transcription. Antisense sequences can be DNA, RNA, or nucleic acid mimics and analogs. (See, e.g., Rossi, J.J. et al. (1991) Antisense Res. Dev. 1(3):285-288; Lee, R. et al. (1998) Biochemistry 37(3):900-1010; Pardridge, W.M. et al. (1995) Proc. Natl. Acad. Sci. USA 92(12):5592-5596; and Nielsen, P. E. and Haaima, G. (1997) Chem. Soc. Rev. 96:73-78.) Typically, the binding which results in modulation of expression occurs through hybridization or binding of complementary base pairs. Antisense sequences can also bind to DNA duplexes through specific interactions in the major groove of the double helix.

The polynucleotides of the present invention and fragments thereof can be used as antisense sequences to modify the expression of the polypeptide encoded by sptm. The antisense sequences can be produced <u>ex vivo</u>, such as by using any of the ABI nucleic acid synthesizer series (PE Biosystems) or other automated systems known in the art. Antisense sequences can also be produced biologically, such as by transforming an appropriate host cell with an expression vector containing the sequence of interest. (See, e.g., Agrawal, <u>supra.</u>)

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In therapeutic use, any gene delivery system suitable for introduction of the antisense sequences into appropriate target cells can be used. Antisense sequences can be delivered intracellularly in the form of an expression plasmid which, upon transcription, produces a sequence complementary to at least a portion of the cellular sequence encoding the target protein. (See, e.g., Slater, J.E., et al. (1998) J. Allergy Clin. Immunol. 102(3):469-475; and Scanlon, K.J., et al. (1995) 9(13):1288-1296.)

Antisense sequences can also be introduced intracellularly through the use of viral vectors, such as retrovirus and adeno-associated virus vectors. (See, e.g., Miller, A.D. (1990) Blood 76:271; Ausubel, F.M. et al. (1995) Current Protocols in Molecular Biology, John Wiley & Sons, New York NY; Uckert, W. and W. Walther (1994) Pharmacol. Ther. 63(3):323-347.) Other gene delivery mechanisms include liposome-derived systems, artificial viral envelopes, and other systems known in the art. (See, e.g., Rossi, J.J. (1995) Br. Med. Bull. 51(1):217-225; Boado, R.J. et al. (1998) J. Pharm. Sci. 87(11):1308-1315; and Morris, M.C. et al. (1997) Nucleic Acids Res. 25(14):2730-2736.)

Expression

In order to express a biologically active SPTM, the nucleotide sequences encoding SPTM or fragments thereof may be inserted into an appropriate expression vector, i.e., a vector which contains the necessary elements for transcriptional and translational control of the inserted coding sequence in a suitable host. Methods which are well known to those skilled in the art may be used to construct expression vectors containing sequences encoding SPTM and appropriate transcriptional and translational control elements. These methods include in vitro recombinant DNA techniques, synthetic

In another embodiment, the toxicity of a test compound is assessed by treating a biological sample containing proteins with the test compound. Proteins that are expressed in the treated biological sample are separated so that the amount of each protein can be quantified. The amount of each protein is compared to the amount of the corresponding protein in an untreated biological sample. A difference in the amount of protein between the two samples is indicative of a toxic response to the test compound in the treated sample. Individual proteins are identified by sequencing the amino acid residues of the individual proteins and comparing these partial sequences to the SPTM encoded by polynucleotides of

In another embodiment, the toxicity of a test compound is assessed by treating a biological sample containing proteins with the test compound. Proteins from the biological sample are incubated 10 with antibodies specific to the SPTM encoded by polynucleotides of the present invention. The amount of protein recognized by the antibodies is quantified. The amount of protein in the treated biological sample is compared with the amount in an untreated biological sample. A difference in the amount of protein between the two samples is indicative of a toxic response to the test compound in the treated

Transcript images may be used to profile sptm expression in distinct tissue types. This process can be used to determine cell signaling activity in a particular tissue type relative to this activity in a different tissue type. Transcript images may be used to generate a profile of sptm expression characteristic of diseased tissue. Transcript images of tissues before and after treatment may be used for diagnostic purposes, to monitor the progression of disease, and to monitor the efficacy of drug treatments for diseases which affect cell signaling activity.

Transcript images of cell lines can be used to assess cell signaling activity and/or to identify cell lines that lack or misregulate this activity. Such cell lines may then be treated with pharmaceutical agents, and a transcript image following treatment may indicate the efficacy of these agents in restoring desired levels of this activity. A similar approach may be used to assess the toxicity of pharmaceutical agents as reflected by undesirable changes in cell signaling activity. Candidate pharmaceutical agents may be evaluated by comparing their associated transcript images with those of pharmaceutical agents of known effectiveness.

Antisense Molecules

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The polynucleotides of the present invention are useful in antisense technology. Antisense 30 technology or therapy relies on the modulation of expression of a target protein through the specific binding of an antisense sequence to a target sequence encoding the target protein or directing its expression. (See, e.g., Agrawal, S., ed. (1996) Antisense Therapeutics, Humana Press Inc., Totawa NJ; Alama, A. et al. (1997) Pharmacol. Res. 36(3):171-178; Crooke, S.T. (1997) Adv. Pharmacol. 35

immunodeficiency (SCID)-X1 disease characterized by X-linked inheritance (Cavazzana-Calvo, M. et al. (2000) Science 288:669-672), severe combined immunodeficiency syndrome associated with an inherited adenosine deaminase (ADA) deficiency (Blaese, R.M. et al. (1995) Science 270:475-480; Bordignon, C. et al. (1995) Science 270:470-475), cystic fibrosis (Zabner, J. et al. (1993) Cell 75:207-216; Crystal, R.G. et al. (1995) Hum. Gene Therapy 6:643-666; Crystal, R.G. et al. (1995) Hum. Gene Therapy 6:667-703), thalassemias, familial hypercholesterolemia, and hemophilia resulting from Factor VIII or Factor IX deficiencies (Crystal, R.G. (1995) Science 270:404-410; Verma, I.M. and Somia, N. (1997) Nature 389:239-242)), (ii) express a conditionally lethal gene product (e.g., in the case of cancers which result from unregulated cell proliferation), or (iii) express a protein which affords protection against intracellular parasites (e.g., against human retroviruses, such as human immunodeficiency virus (HIV) (Baltimore, D. (1988) Nature 335:395-396; Poeschla, E. et al. (1996) Proc. Natl. Acad. Sci. USA. 93:11395-11399), hepatitis B or C virus (HBV, HCV); fungal parasites, such as Candida albicans and Paracoccidioides brasiliensis; and protozoan parasites such as <u>Plasmodium falciparum</u> and <u>Trypanosoma cruzi</u>). In the case where a genetic deficiency in sptm expression or regulation causes disease, the expression of sptm from an appropriate population of 15 transduced cells may alleviate the clinical manifestations caused by the genetic deficiency.

In a further embodiment of the invention, diseases or disorders caused by deficiencies in sptm are treated by constructing mammalian expression vectors comprising sptm and introducing these vectors by mechanical means into sptm-deficient cells. Mechanical transfer technologies for use with cells in vivo or ex vitro include (i) direct DNA microinjection into individual cells, (ii) ballistic gold particle delivery, (iii) liposome-mediated transfection, (iv) receptor-mediated gene transfer, and (v) the use of DNA transposons (Morgan, R.A. and Anderson, W.F. (1993) Annu. Rev. Biochem. 62:191-217; Ivics, Z. (1997) Cell 91:501-510; Boulay, J-L. and Récipon, H. (1998) Curr. Opin. Biotechnol. 9:445-450).

Expression vectors that may be effective for the expression of sptm include, but are not limited to, the PCDNA 3.1, EPITAG, PRCCMV2, PREP, PVAX vectors (Invitrogen, Carlsbad CA), PCMV-SCRIPT, PCMV-TAG, PEGSH/PERV (Stratagene, La Jolla CA), and PTET-OFF, PTET-ON, PTRE2, PTRE2-LUC, PTK-HYG (Clontech, Palo Alto CA). The sptm of the invention may be expressed using (i) a constitutively active promoter, (e.g., from cytomegalovirus (CMV), Rous sarcoma virus (RSV), SV40 virus, thymidine kinase (TK), or β-actin genes), (ii) an inducible promoter (e.g., the tetracycline-regulated promoter (Gossen, M. and Bujard, H. (1992) Proc. Natl. Acad. Sci. U.S.A. 89:5547-5551; Gossen, M. et al., (1995) Science 268:1766-1769; Rossi, F.M.V. and Blau, H.M. (1998) Curr. Opin. Biotechnol. 9:451-456), commercially available in the T-REX plasmid (Invitrogen); the ecdysone-inducible promoter (available in the plasmids PVGRXR and PIND;

Invitrogen); the FK506/rapamycin inducible promoter; or the RU486/mifepristone inducible promoter

techniques, and in vivo genetic recombination. (See, e.g., Sambrook, supra, Chapters 4, 8, 16, and 17; and Ausubel, supra, Chapters 9, 10, 13, and 16.)

A variety of expression vector/host systems may be utilized to contain and express sequences encoding SPTM. These include, but are not limited to, microorganisms such as bacteria transformed with recombinant bacteriophage, plasmid, or cosmid DNA expression vectors; yeast transformed with yeast expression vectors; insect cell systems infected with viral expression vectors (e.g., baculovirus); plant cell systems transformed with viral expression vectors (e.g., cauliflower mosaic virus, CaMV, or tobacco mosaic virus, TMV) or with bacterial expression vectors (e.g., Ti or pBR322 plasmids); or animal (mammalian) cell systems. (See, e.g., Sambrook, supra; Ausubel, 1995, supra, Van Heeke, G. and S.M. Schuster (1989) J. Biol. Chem. 264:5503-5509; Bitter, G.A. et al. (1987) Methods Enzymol. 10 153:516-544; Scorer, C.A. et al. (1994) Bio/Technology 12:181-184; Engelhard, E.K. et al. (1994) Proc. Natl. Acad. Sci. USA 91:3224-3227; Sandig, V. et al. (1996) Hum. Gene Ther. 7:1937-1945; Takamatsu, N. (1987) EMBO J. 6:307-311; Coruzzi, G. et al. (1984) EMBO J. 3:1671-1680; Broglie, R. et al. (1984) Science 224:838-843; Winter, J. et al. (1991) Results Probl. Cell Differ. 17:85-105; The McGraw Hill Yearbook of Science and Technology (1992) McGraw Hill, New York NY, pp. 15 191-196; Logan, J. and T. Shenk (1984) Proc. Natl. Acad. Sci. USA 81:3655-3659; and Harrington, J.J. et al. (1997) Nat. Genet. 15:345-355.) Expression vectors derived from retroviruses, adenoviruses, or herpes or vaccinia viruses, or from various bacterial plasmids, may be used for delivery of nucleotide sequences to the targeted organ, tissue, or cell population. (See, e.g., Di Nicola, M. et al. (1998) Cancer Gen. Ther. 5(6):350-356; Yu, M. et al., (1993) Proc. Natl. Acad. Sci. USA 90(13):6340-6344; Buller, R.M. et al. (1985) Nature 317(6040):813-815; McGregor, D.P. et al. (1994) Mol. Immunol.

limited by the host cell employed. For long term production of recombinant proteins in mammalian systems, stable expression of SPTM in cell lines is preferred. For example, sequences encoding SPTM can be transformed into cell lines using expression vectors which may contain viral origins of replication and/or endogenous expression elements and a selectable marker gene on the same or on a separate vector. Any number of selection systems may be used to recover transformed cell lines. (See, e.g., Wigler, M. et al. (1977) Cell 11:223-232; Lowy, I. et al. (1980) Cell 22:817-823.; Wigler, M. et al. (1980) Proc. Natl. Acad. Sci. USA 77:3567-3570; Colbere-Garapin, F. et al. (1981) J. Mol. Biol. 150:1-14; Hartman, S.C. and 30 R.C.Mulligan (1988) Proc. Natl. Acad. Sci. USA 85:8047-8051; Rhodes, C.A. (1995) Methods Mol.

31(3):219-226; and Verma, I.M. and N. Somia (1997) Nature 389:239-242.) The invention is not

Therapeutic Uses of sptm

The polynucleotides encoding SPTM may be used for somatic or germline gene therapy. Gene therapy may be performed to (i) correct a genetic deficiency (e.g., in the cases of severe combined 35

Number 5,707,618 to Armentano ("Adenovirus vectors for gene therapy"), hereby incorporated by reference. For adenoviral vectors, see also Antinozzi, P.A. et al. (1999) Annu. Rev. Nutr. 19:511-544 and Verma, I.M. and Somia, N. (1997) Nature 18:389:239-242, both incorporated by reference herein.

In another alternative, a herpes-based, gene therapy delivery system is used to deliver sptm to target cells which have one or more genetic abnormalities with respect to the expression of sptm. The use of herpes simplex virus (HSV)-based vectors may be especially valuable for introducing sptm to cells of the central nervous system, for which HSV has a tropism. The construction and packaging of herpes-based vectors are well known to those with ordinary skill in the art. A replication-competent herpes simplex virus (HSV) type 1-based vector has been used to deliver a reporter gene to the eyes of primates (Liu, X. et al. (1999) Exp. Eye Res. 169:385-395). The construction of a HSV-1 virus vector has also been disclosed in detail in U.S. Patent Number 5,804,413 to DeLuca ("Herpes simplex virus strains for gene transfer"), which is hereby incorporated by reference. U.S. Patent Number 5,804,413 teaches the use of recombinant HSV d92 which consists of a genome containing at least one exogenous gene to be transferred to a cell under the control of the appropriate promoter for purposes including human gene therapy. Also taught by this patent are the construction and use of recombinant HSV.... strains deleted for ICP4, ICP27 and ICP22. For HSV vectors, see also Goins, W. F. et al. 1999 J. Virol. 73:519-532 and Xu, H. et al., (1994) Dev. Biol. 163:152-161, hereby incorporated by reference. The manipulation of cloned herpesvirus sequences, the generation of recombinant virus following the transfection of multiple plasmids containing different segments of the large herpesvirus genomes, the growth and propagation of herpesvirus, and the infection of cells with herpesvirus are techniques well known to those of ordinary skill in the art.

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In another alternative, an alphavirus (positive, single-stranded RNA virus) vector is used to deliver sptm to target cells. The biology of the prototypic alphavirus, Semliki Forest Virus (SFV), has been studied extensively and gene transfer vectors have been based on the SFV genome (Garoff, H. and Li, K-J. (1998) Curr. Opin. Biotech. 9:464-469). During alphavirus RNA replication, a subgenomic RNA is generated that normally encodes the viral capsid proteins. This subgenomic RNA replicates to higher levels than the full-length genomic RNA, resulting in the overproduction of capsid proteins relative to the viral proteins with enzymatic activity (e.g., protease and polymerase). Similarly, inserting sptm into the alphavirus genome in place of the capsid-coding region results in the production of a large number of sptm RNAs and the synthesis of high levels of SPTM in vector transduced cells. While alphavirus infection is typically associated with cell lysis within a few days, the ability to establish a persistent infection in hamster normal kidney cells (BHK-21) with a variant of Sindbis virus (SIN) indicates that the lytic replication of alphaviruses can be altered to suit the needs of the gene therapy application (Dryga, S.A. et al. (1997) Virology 228:74-83). The wide host range of alphaviruses will allow the introduction of sptm into a variety of cell types. The specific transduction

(Rossi, F.M.V. and Blau, H.M. supra), or (iii) a tissue-specific promoter or the native promoter of the endogenous gene encoding SPTM from a normal individual.

Commercially available liposome transformation kits (e.g., the PERFECT LIPID TRANSFECTION KIT, available from Invitrogen) allow one with ordinary skill in the art to deliver polynucleotides to target cells in culture and require minimal effort to optimize experimental parameters. In the alternative, transformation is performed using the calcium phosphate method (Graham, F.L. and Eb, A.J. (1973) Virology 52:456-467), or by electroporation (Neumann, E. et al. (1982) EMBO J. 1:841-845). The introduction of DNA to primary cells requires modification of these standardized mammalian transfection protocols.

In another embodiment of the invention, diseases or disorders caused by genetic defects with 10 respect to sptm expression are treated by constructing a retrovirus vector consisting of (i) sptm under the control of an independent promoter or the retrovirus long terminal repeat (LTR) promoter, (ii) appropriate RNA packaging signals, and (iii) a Rev-responsive element (RRE) along with additional retrovirus cis-acting RNA sequences and coding sequences required for efficient vector propagation. Retrovirus vectors (e.g., PFB and PFBNEO) are commercially available (Stratagene) and are based on published data (Riviere, I. et al. (1995) Proc. Natl. Acad. Sci. U.S.A. 92:6733-6737), incorporated by reference herein. The vector is propagated in an appropriate vector producing cell line (VPCL) that expresses an envelope gene with a tropism for receptors on the target cells or a promiscuous envelope protein such as VSVg (Armentano, D. et al. (1987) J. Virol. 61:1647-1650; Bender, M.A. et al. (1987) J. Virol. 61:1639-1646; Adam, M.A. and Miller, A.D. (1988) J. Virol. 62:3802-3806; Dull, T. et al. (1998) J. Virol. 72:8463-8471; Zufferey, R. et al. (1998) J. Virol. 72:9873-9880). U.S. Patent Number 5,910,434 to Rigg ("Method for obtaining retrovirus packaging cell lines producing high transducing efficiency retroviral supernatant") discloses a method for obtaining retrovirus packaging cell lines and is hereby incorporated by reference. Propagation of retrovirus vectors, transduction of a population of cells (e.g., CD4+ T-cells), and the return of transduced cells to a patient are procedures well known to persons skilled in the art of gene therapy and have been well documented (Ranga, U. et al. (1997) J. Virol. 71:7020-7029; Bauer, G. et al. (1997) Blood 89:2259-2267; Bonyhadi, M.L. (1997) J. Virol. 71:4707-4716; Ranga, U. et al. (1998) Proc. Natl. Acad. Sci. U.S.A. 95:1201-1206; Su, L. (1997) Blood 89:2283-2290). 30

In the alternative, an adenovirus-based gene therapy delivery system is used to deliver sptm to cells which have one or more genetic abnormalities with respect to the expression of sptm. The construction and packaging of adenovirus-based vectors are well known to those with ordinary skill in the art. Replication defective adenovirus vectors have proven to be versatile for importing genes encoding immunoregulatory proteins into intact islets in the pancreas (Csete, M.E. et al. (1995)

cells are then used to produce hybridomas using standard techniques. About 20 mg of peptide is sufficient for labeling and screening several thousand clones. Hybridomas of interest are detected by screening with radioiodinated peptide to identify those fusions producing peptide-specific monoclonal antibody. In a typical protocol, wells of a multi-well plate (FAST, Becton-Dickinson, Palo Alto, CA) are coated with affinity-purified, specific rabbit-anti-mouse (or suitable anti-species IgG) antibodies at 10 mg/ml. The coated wells are blocked with 1% BSA and washed and exposed to supernatants from hybridomas. After incubation, the wells are exposed to radiolabeled peptide at 1 mg/ml.

Clones producing antibodies bind a quantity of labeled peptide that is detectable above background. Such clones are expanded and subjected to 2 cycles of cloning. Cloned hybridomas are injected into pristane-treated mice to produce ascites, and monoclonal antibody is purified from the ascitic fluid by affinity chromatography on protein A (Amersham Pharmacia Biotech). Several procedures for the production of monoclonal antibodies, including in vitro production, are described in Pound (supra). Monoclonal antibodies with antipeptide activity are tested for anti-SPTM activity using protocols well known in the art, including ELISA, RIA, and immunoblotting.

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Antibody fragments containing specific binding sites for an epitope may also be generated. For example, such fragments include, but are not limited to, the F(ab')2 fragments produced by pepsin digestion of the antibody molecule, and the Fab fragments generated by reducing the disulfide bridges of the F(ab')2 fragments. Alternatively, construction of Fab expression libraries in filamentous bacteriophage allows rapid and easy identification of monoclonal fragments with desired specificity (Pound, supra, Chaps. 45-47). Antibodies generated against polypeptide encoded by sptm can be used to purify and characterize full-length SPTM protein and its activity, binding partners, etc.

Assays Using Antibodies

Anti-SPTM antibodies may be used in assays to quantify the amount of SPTM found in a particular human cell. Such assays include methods utilizing the antibody and a label to detect expression level under normal or disease conditions. The peptides and antibodies of the invention may be used with or without modification or labeled by joining them, either covalently or noncovalently, with a reporter molecule.

Protocols for detecting and measuring protein expression using either polyclonal or monoclonal antibodies are well known in the art. Examples include ELISA, RIA, and fluorescent activated cell sorting (FACS). Such immunoassays typically involve the formation of complexes between the SPTM and its specific antibody and the measurement of such complexes. These and other assays are described in Pound (supra).

Without further elaboration, it is believed that one skilled in the art can, using the preceding description, utilize the present invention to its fullest extent. The following preferred specific

of a subset of cells in a population may require the sorting of cells prior to transduction. The methods of manipulating infectious cDNA clones of alphaviruses, performing alphavirus cDNA and RNA transfections, and performing alphavirus infections, are well known to those with ordinary skill in the

Antibodies

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Anti-SPTM antibodies may be used to analyze protein expression levels. Such antibodies include, but are not limited to, polyclonal, monoclonal, chimeric, single chain, and Fab fragments. For descriptions of and protocols of antibody technologies, see, e.g., Pound J.D. (1998) Immunochemical Protocols, Humana Press, Totowa, NJ.

The amino acid sequence encoded by the sptm of the Sequence Listing may be analyzed by appropriate software (e.g., LASERGENE NAVIGATOR software, DNASTAR) to determine regions of high immunogenicity. The optimal sequences for immunization are selected from the C-terminus, the N-terminus, and those intervening, hydrophilic regions of the polypeptide which are likely to be exposed to the external environment when the polypeptide is in its natural conformation. Analysis used to select appropriate epitopes is also described by Ausubel (1997, supra, Chapter 11.7). Peptides used for antibody induction do not need to have biological activity; however, they must be antigenic. Peptides used to induce specific antibodies may have an amino acid sequence consisting of at five amino acids, preferably at least 10 amino acids, and most preferably 15 amino acids. A peptide which mimics an antigenic fragment of the natural polypeptide may be fused with another protein such as keyhole limpet cyanin (KLH; Sigma, St. Louis MO) for antibody production. A peptide encompassing an antigenic region may be expressed from an sptm, synthesized as described above, or purified from human cells.

Procedures well known in the art may be used for the production of antibodies. Various hosts including mice, goats, and rabbits, may be immunized by injection with a peptide. Depending on the host species, various adjuvants may be used to increase immunological response.

25 In one procedure, peptides about 15 residues in length may be synthesized using an ABI 431A peptide synthesizer (PE Biosystems) using fmoc-chemistry and coupled to KLH (Sigma) by reaction with M-maleimidobenzoyl-N-hydroxysuccinimide ester (Ausubel, 1995, supra). Rabbits are immunized with the peptide-KLH complex in complete Freund's adjuvant. The resulting antisera are tested for antipeptide activity by binding the peptide to plastic, blocking with 1% bovine serum albumin (BSA), reacting with rabbit antisera, washing, and reacting with radioiodinated goat anti-rabbit IgG. Antisera with antipeptide activity are tested for anti-SPTM activity using protocols well known in the art, including ELISA, radioimmunoassay (RIA), and immunoblotting.

In another procedure, isolated and purified peptide may be used to immunize mice (about 100 μg of peptide) or rabbits (about 1 mg of peptide). Subsequently, the peptide is radioiodinated and used to screen the immunized animals' B-lymphocytes for production of antipeptide antibodies. Positive

Plasmids were recovered from host cells by <u>in vivo</u> excision using the UNIZAP vector system (Stratagene) or by cell lysis. Plasmids were purified using at least one of the following: the Magic or WIZARD Minipreps DNA purification system (Promega); the AGTC Miniprep purification kit (Edge BioSystems, Gaithersburg MD); and the QIAWELL 8, QIAWELL 8 Plus, and QIAWELL 8 Ultra plasmid purification systems or the R.E.A.L. PREP 96 plasmid purification kit (QIAGEN). Following precipitation, plasmids were resuspended in 0.1 ml of distilled water and stored, with or without lyophilization, at 4°C.

Alternatively, plasmid DNA was amplified from host cell lysates using direct link PCR in a high-throughput format. (Rao, V.B. (1994) Anal. Biochem. 216:1-14.) Host cell lysis and thermal cycling steps were carried out in a single reaction mixture. Samples were processed and stored in 384-well plates, and the concentration of amplified plasmid DNA was quantified fluorometrically using PICOGREEN dye (Molecular Probes, Inc. (Molecular Probes), Eugene OR) and a FLUOROSKAN II fluorescence scanner (Labsystems Oy, Helsinki, Finland).

III. Sequencing and Analysis

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cDNA sequencing reactions were processed using standard methods or high-throughput instrumentation such as the ABI CATALYST 800 thermal cycler (PE Biosystems) or the PTC-200 thermal cycler (MJ Research) in conjunction with the HYDRA microdispenser (Robbins Scientific Corp., Sunnyvale CA) or the MICROLAB 2200 liquid transfer system (Hamilton). cDNA sequencing reactions were prepared using reagents provided by Amersham Pharmacia Biotech or supplied in ABI sequencing kits such as the ABI PRISM BIGDYE Terminator cycle sequencing ready reaction kit (PE Biosystems). Electrophoretic separation of cDNA sequencing reactions and detection of labeled polynucleotides were carried out using the MEGABACE 1000 DNA sequencing system (Molecular Dynamics); the ABI PRISM 373 or 377 sequencing system (PE Biosystems) in conjunction with standard ABI protocols and base calling software; or other sequence analysis systems known in the art. Reading frames within the cDNA sequences were identified using standard methods (reviewed in Ausubel, 1997, supra, Chapter 7.7). Some of the cDNA sequences were selected for extension using the techniques disclosed in Example VIII.

IV. Assembly and Analysis of Sequences

Component sequences from chromatograms were subject to PHRED analysis and assigned a quality score. The sequences having at least a required quality score were subject to various pre-processing editing pathways to eliminate, e.g., low quality 3' ends, vector and linker sequences, polyA tails, Alu repeats, mitochondrial and ribosomal sequences, bacterial contamination sequences, and sequences smaller than 50 base pairs. In particular, low-information sequences and repetitive elements (e.g., dinucleotide repeats, Alu repeats, etc.) were replaced by "n's", or masked, to prevent spurious matches.

embodiments are, therefore, to be construed as merely illustrative, and not limitative of the remainder of the disclosure in any way whatsoever.

The disclosures of all patents, applications, and publications mentioned above and below, in particular U.S. Ser. No. 60/156,624, U.S. Ser. No. 60/156,625, U.S. Ser. No. 60/168,614, U.S. Ser. No. 60/168,611, and U.S. Ser. No. 60/168,613 are hereby expressly incorporated by reference.

EXAMPLES

I. Construction of cDNA Libraries

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RNA was purchased from CLONTECH Laboratories, Inc. (Palo Alto CA) or isolated from various tissues. Some tissues were homogenized and lysed in guanidinium isothiocyanate, while others were homogenized and lysed in phenol or in a suitable mixture of denaturants, such as TRIZOL (Life Technologies), a monophasic solution of phenol and guanidine isothiocyanate. The resulting lysates were centrifuged over CsCl cushions or extracted with chloroform. RNA was precipitated with either isopropanol or sodium acetate and ethanol, or by other routine methods.

Phenol extraction and precipitation of RNA were repeated as necessary to increase RNA. purity. In most cases, RNA was treated with DNase. For most libraries, poly(A+) RNA was isolated using oligo d(T)-coupled paramagnetic particles (Promega Corporation (Promega), Madison WI), OLIGOTEX latex particles (QIAGEN, Inc. (QIAGEN), Valencia CA), or an OLIGOTEX mRNA purification kit (QIAGEN). Alternatively, RNA was isolated directly from tissue lysates using other RNA isolation kits, e.g., the POLY(A)PURE mRNA purification kit (Ambion, Inc., Austin TX).

In some cases, Stratagene was provided with RNA and constructed the corresponding cDNA 20 libraries. Otherwise, cDNA was synthesized and cDNA libraries were constructed with the UNIZAP vector system (Stratagene Cloning Systems, Inc. (Stratagene), La Jolla CA) or SUPERSCRIPT plasmid system (Life Technologies), using the recommended procedures or similar methods known in the art. (See, e.g., Ausubel, 1997, supra, Chapters 5.1 through 6.6.) Reverse transcription was initiated using oligo d(T) or random primers. Synthetic oligonucleotide adapters were ligated to double stranded cDNA, and the cDNA was digested with the appropriate restriction enzyme or enzymes. For most libraries, the cDNA was size-selected (300-1000 bp) using SEPHACRYL S1000, SEPHAROSE CL2B, or SEPHAROSE CL4B column chromatography (Amersham Pharmacia Biotech) or preparative agarose gel electrophoresis. cDNAs were ligated into compatible restriction enzyme sites of the polylinker of a suitable plasmid, e.g., PBLUESCRIPT plasmid (Stratagene), pSPORT1 plasmid (Life Technologies), or pINCY (Incyte). Recombinant plasmids were transformed into competent <u>E.</u> coli cells including XL1-Blue, XL1-BlueMRF, or SOLR from Stratagene or DH5\alpha, DH10B, or ElectroMAX DH10B from Life Technologies.

II. Isolati n of cDNA Clones

Following assembly, template sequences were subjected to motif, BLAST, and functional analyses, and categorized in protein hierarchies using methods described in, e.g., "Database System Employing Protein Function Hierarchies for Viewing Biomolecular Sequence Data," U.S.S.N. 08/812,290, filed March 6, 1997; "Relational Database for Storing Biomolecule Information," U.S.S.N. 08/947,845, filed October 9, 1997; "Project-Based Full-Length Biomolecular Sequence Database," U.S.S.N. 08/811,758, filed March 6, 1997; and "Relational Database and System for Storing Information Relating to Biomolecular Sequences," U.S.S.N. 09/034,807, filed March 4, 1998, all of which are incorporated by reference herein.

The template sequences were further analyzed by translating each template in all three forward reading frames and searching each translation against the Pfam database of hidden Markov model-based protein families and domains using the HMMER software package (available to the public from Washington University School of Medicine, St. Louis MO). (See also World Wide Web site http://pfam.wustl.edu/ for detailed descriptions of Pfam protein domains and families.)

Additionally, the template sequences were translated in all three forward reading frames, and each translation was searched against hidden Markov models for signal peptide and transmembrane domains using the HMMER software package. Construction of hidden Markov models and their usage in sequence analysis has been described. (See, for example, Eddy, S.R. (1996) Curr. Opin. Str. Biol. 6:361-365.) Regions of templates which, when translated, contain similarity to signal peptide or transmembrane domain consensus sequences are reported in Table 1. Only those signal peptide or transmembrane hits with a cutoff score of 11 bits or greater are reported. A cutoff score of 11 bits or greater corresponds to at least about 91-94% true-positives in signal peptide prediction, and at least about 75% true-positives in transmembrane domain prediction.

Template sequences are further analyzed using the bioinformatics tools listed in Table 4, or using sequence analysis software known in the art such as MACDNASIS PRO software (Hitachi Software Engineering, South San Francisco CA) and LASERGENE software (DNASTAR). Template sequences may be further queried against public databases such as the GenBank rodent, mammalian, vertebrate, prokaryote, and eukaryote databases.

V. Analysis of Polynucleotide Expression

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Northern analysis is a laboratory technique used to detect the presence of a transcript of a gene and involves the hybridization of a labeled nucleotide sequence to a membrane on which RNAs from a particular cell type or tissue have been bound. (See, e.g., Sambrook, supra, ch. 7; Ausubel, 1995, supra, ch. 4 and 16.)

Analogous computer techniques applying BLAST were used to search for identical or related molecules in cDNA databases such as GenBank or LIFESEQ (Incyte Genomics). This analysis is much faster than multiple membrane-based hybridizations. In addition, the sensitivity of the computer

Processed sequences were then subject to assembly procedures in which the sequences were assigned to gene bins (bins). Each sequence could only belong to one bin. Sequences in each gene bin were assembled to produce consensus sequences (templates). Subsequent new sequences were added to existing bins using BLASTn (v.1.4 WashU) and CROSSMATCH. Candidate pairs were identified as all BLAST hits having a quality score greater than or equal to 150. Alignments of at least 82% local identity were accepted into the bin. The component sequences from each bin were assembled using a version of PHRAP. Bins with several overlapping component sequences were assembled using DEEP PHRAP. The orientation (sense or antisense) of each assembled template was determined based on the number and orientation of its component sequences. Template sequences as disclosed in the sequence listing correspond to sense strand sequences (the "forward" reading frames), to the best determination. The complementary (antisense) strands are inherently disclosed herein. The component sequences which were used to assemble each template consensus sequence are listed in Table 2, along with their positions along the template nucleotide sequences.

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Bins were compared against each other and those having local similarity of at least 82% were combined and reassembled. Reassembled bins having templates of insufficient overlap (less than 95% local identity) were re-split. Assembled templates were also subject to analysis by STITCHER/EXON MAPPER algorithms which analyze the probabilities of the presence of splice variants, alternatively spliced exons, splice junctions, differential expression of alternative spliced genes across tissue types or disease states, etc. These resulting bins were subject to several rounds of the above assembly procedures.

Once gene bins were generated based upon sequence alignments, bins were clone joined based upon clone information. If the 5' sequence of one clone was present in one bin and the 3' sequence from the same clone was present in a different bin, it was likely that the two bins actually belonged together in a single bin. The resulting combined bins underwent assembly procedures to regenerate the consensus sequences.

The final assembled templates were subsequently annotated using the following procedure. Template sequences were analyzed using BLASTn (v2.0, NCBI) versus gbpri (GenBank version 118). "Hits" were defined as an exact match having from 95% local identity over 200 base pairs through 100% local identity over 100 base pairs, or a homolog match having an E-value, i.e. a probability score, of $\leq 1 \times 10^{-8}$. The hits were subject to frameshift FASTx versus GENPEPT (GenBank version 118). (See Table 4). In this analysis, a homolog match was defined as having an E-value of $\leq 1 \times 10^{-8}$. The assembly method used above was described in "System and Methods for Analyzing Biomolecular Sequences," U.S.S.N. 09/276,534, filed March 25, 1999, and the LIFESEQ Gold user manual (Incyte) both incorporated by reference herein.

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sequences and cDNA library/tissue information are found in the LIFESEQ GOLD database (Incyte Genomics, Palo Alto CA).

Tissue Distribution Profiling

A tissue distribution profile is determined for each template by compiling the cDNA library VI. tissue classifications of its component cDNA sequences. Each component sequence, is derived from a cDNA library constructed from a human tissue. Each human tissue is classified into one of the following categories: cardiovascular system; connective tissue; digestive system; embryonic structures; endocrine system; exocrine glands; genitalia, female; genitalia, male; germ cells; hemic and immune system; liver; musculoskeletal system; nervous system; pancreas; respiratory system; sense organs; skin; stomatognathic system; unclassified/mixed; or urinary tract. Template sequences, component sequences, and cDNA library/tissue information are found in the LIFESEQ GOLD database (Incyte 10 Genomics, Palo Alto CA).

Table 3 shows the tissue distribution profile for the templates of the invention. For each template, the three most frequently observed tissue categories are shown in column 3, along with the percentage of component sequences belonging to each category. Only tissue categories with percentage values of ≥10% are shown. A tissue distribution of "widely distributed" in column 3 = indicates percentage values of <10% in all tissue categories.

Transcript Image Analysis VII.

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Transcript images are generated as described in Seilhamer et al., "Comparative Gene Transcript Analysis," U.S. Patent Number 5,840,484, incorporated herein by reference.

VIII. Extension of Polynucleotide Sequences and Isolation of a Full-length cDNA

Oligonucleotide primers designed using an sptm of the Sequence Listing are used to extend the nucleic acid sequence. One primer is synthesized to initiate 5' extension of the template, and the other primer, to initiate 3' extension of the template. The initial primers may be designed using OLIGO 4.06 software (National Biosciences, Inc. (National Biosciences), Plymouth MN), or another appropriate program, to be about 22 to 30 nucleotides in length, to have a GC content of about 50% or more, and to anneal to the target sequence at temperatures of about 68°C to about 72°C. Any stretch of nucleotides which would result in hairpin structures and primer-primer dimerizations are avoided. Selected human cDNA libraries are used to extend the sequence. If more than one extension is necessary or desired, additional or nested sets of primers are designed.

High fidelity amplification is obtained by PCR using methods well known in the art. PCR is performed in 96-well plates using the PTC-200 thermal cycler (MJ Research). The reaction mix contains DNA template, 200 nmol of each primer, reaction buffer containing Mg²⁺, (NH₄)₂SO₄, and βmercaptoethanol, Taq DNA polymerase (Amersham Pharmacia Biotech), ELONGASE enzyme (Life Technologies), and Pfu DNA polymerase (Stratagene), with the following parameters for primer pair

search can be modified to determine whether any particular match is categorized as exact or similar. The basis of the search is the product score, which is defined as:

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BLAST Score x Percent Identity

5 x minimum {length(Seq. 1), length(Seq. 2)}

The product score takes into account both the degree of similarity between two sequences and the length of the sequence match. The product score is a normalized value between 0 and 100, and is calculated as follows: the BLAST score is multiplied by the percent nucleotide identity and the product is divided by (5 times the length of the shorter of the two sequences). The BLAST score is calculated by assigning a score of +5 for every base that matches in a high-scoring segment pair (HSP), and -4 for every mismatch. Two sequences may share more than one HSP (separated by gaps). If there is more than one HSP, then the pair with the highest BLAST score is used to calculate the product score. The product score represents a balance between fractional overlap and quality in a BLAST alignment. For example, a product score of 100 is produced only for 100% identity over the entire length of the shorter of the two sequences being compared. A product score of 70 is produced either by 100% identity and 70% overlap at one end, or by 88% identity and 100% overlap at the other. A product score of 50 is produced either by 100% identity and 50% overlap at one end, or 79% identity and 100% overlap.

Alternatively, polynucleotide sequences encoding SPTM are analyzed with respect to the tissue sources from which they were derived. Polynucleotide sequences encoding SPTM were assembled, at 20 least in part, with overlapping Incyte cDNA sequences. Each cDNA sequence is derived from a cDNA library constructed from a human tissue. Each human tissue is classified into one of the following organ/tissue categories: cardiovascular system; connective tissue; digestive system; embryonic structures; endocrine system; exocrine glands; genitalia, female; genitalia, male; germ cells; hemic and immune system; liver; musculoskeletal system; nervous system; pancreas; respiratory system; sense organs; skin; stomatognathic system; unclassified/mixed; or urinary tract. The number of libraries in each category for each polynucleotide sequence encoding SPTM is counted and divided by the total number of libraries across all categories for each polynucleotide sequence encoding SPTM. Similarly, each human tissue is classified into one of the following disease/condition categories: cancer, cell line, developmental, inflammation, neurological, trauma, cardiovascular, pooled, and other, and the number 30 of libraries in each category for each polynucleotide sequence encoding SPTM is counted and divided by the total number of libraries across all categories for each polynucleotide sequence encoding SPTM. The resulting percentages reflect the tissue- and disease-specific expression of cDNA encoding SPTM. Percentage values of tissue-specific and disease-specific expression are reported in Table 3. cDNA

Hybridization probes derived from the sptm of the Sequence Listing are employed for screening cDNAs, mRNAs, or genomic DNA. The labeling of probe nucleotides between 100 and 1000 nucleotides in length is specifically described, but essentially the same procedure may be used with larger cDNA fragments. Probe sequences are labeled at room temperature for 30 minutes using a T4 polynucleotide kinase, γ^{32} P-ATP, and 0.5X One-Phor-All Plus (Amersham Pharmacia Biotech) buffer and purified using a ProbeQuant G-50 Microcolumn (Amersham Pharmacia Biotech). The probe mixture is diluted to 10^7 dpm/ μ g/ml hybridization buffer and used in a typical membrane-based hybridization analysis.

The DNA is digested with a restriction endonuclease such as Eco RV and is electrophoresed through a 0.7% agarose gel. The DNA fragments are transferred from the agarose to nylon membrane (NYTRAN Plus, Schleicher & Schuell, Inc., Keene NH) using procedures specified by the manufacturer of the membrane. Prehybridization is carried out for three or more hours at 68°C, and hybridization is carried out overnight at 68°C. To remove non-specific signals, blots are sequentially washed at room temperature under increasingly stringent conditions, up to 0.1x saline sodium citrate (SSC) and 0.5% sodium dodecyl sulfate. After the blots are placed in a PHOSPHORIMAGER cassette (Molecular Dynamics) or are exposed to autoradiography film, hybridization patterns of standard and experimental lanes are compared. Essentially the same procedure is employed when screening RNA.

X. Chromosome Mapping of sptm

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The cDNA sequences which were used to assemble SEQ ID NO:1-63 are compared waiting sequences from the Incyte LIFESEQ database and public domain databases using BLAST and other implementations of the Smith-Waterman algorithm. Sequences from these databases that match SEO ID NO:1-63 are assembled into clusters of contiguous and overlapping sequences using assembly algorithms such as PHRAP (Table 4). Radiation hybrid and genetic mapping data available from public resources such as the Stanford Human Genome Center (SHGC), Whitehead Institute for Genome Research (WIGR), and Généthon are used to determine if any of the clustered sequences have been previously mapped. Inclusion of a mapped sequence in a cluster will result in the assignment of all sequences of that cluster, including its particular SEQ ID NO:, to that map location. The genetic map locations of SEQ ID NO:1-63 are described as ranges, or intervals, of human chromosomes. The map position of an interval, in centiMorgans, is measured relative to the terminus of the chromosome's parm. (The centiMorgan (cM) is a unit of measurement based on recombination frequencies between chromosomal markers. On average, 1 cM is roughly equivalent to 1 megabase (Mb) of DNA in humans, although this can vary widely due to hot and cold spots of recombination.) The cM distances are based on genetic markers mapped by Généthon which provide boundaries for radiation hybrid markers whose sequences were included in each of the clusters.

35 XI. Micr array Analysis

PCI A and PCI B: Step 1: 94°C, 3 min; Step 2: 94°C, 15 sec; Step 3: 60°C, 1 min; Step 4: 68°C, 2 min; Step 5: Steps 2, 3, and 4 repeated 20 times; Step 6: 68°C, 5 min; Step 7: storage at 4°C. In the alternative, the parameters for primer pair T7 and SK+ are as follows: Step 1: 94°C, 3 min; Step 2: 94°C, 15 sec; Step 3: 57°C, 1 min; Step 4: 68°C, 2 min; Step 5: Steps 2, 3, and 4 repeated 20 times; Step 6: 68°C, 5 min; Step 7: storage at 4°C.

The concentration of DNA in each well is determined by dispensing $100~\mu l$ PICOGREEN quantitation reagent (0.25% (v/v); Molecular Probes) dissolved in 1X Tris-EDTA (TE) and 0.5 μl of undiluted PCR product into each well of an opaque fluorimeter plate (Corning Incorporated (Corning), Corning NY), allowing the DNA to bind to the reagent. The plate is scanned in a FLUOROSKAN II (Labsystems Oy) to measure the fluorescence of the sample and to quantify the concentration of DNA A 5 μl to 10 μl aliquot of the reaction mixture is analyzed by electrophoresis on a 1% agarose mini-gel to determine which reactions are successful in extending the sequence.

The extended nucleotides are desalted and concentrated, transferred to 384-well plates, digested with CviJI cholera virus endonuclease (Molecular Biology Research, Madison WI), and sonicated or sheared prior to religation into pUC 18 vector (Amersham Pharmacia Biotech). For shotgun sequencing, the digested nucleotides are separated on low concentration (0.6 to 0.8%) agarose gels, fragments are excised, and agar digested with AGAR ACE (Promega). Extended clones are religated using T4 ligase (New England Biolabs, Inc., Beverly MA) into pUC 18 vector (Amersham Pharmacia Biotech), treated with Pfu DNA polymerase (Stratagene) to fill-in restriction site overhangs, and transfected into competent <u>E. coli</u> cells. Transformed cells are selected on antibiotic-containing media, individual colonies are picked and cultured overnight at 37°C in 384-well plates in LB/2x carbenicillin liquid media.

The cells are lysed, and DNA is amplified by PCR using Taq DNA polymerase (Amersham Pharmacia Biotech) and Pfu DNA polymerase (Stratagene) with the following parameters: Step 1: 94°C, 3 min; Step 2: 94°C, 15 sec; Step 3: 60°C, 1 min; Step 4: 72°C, 2 min; Step 5: steps 2, 3, and 4 repeated 29 times; Step 6: 72°C, 5 min; Step 7: storage at 4°C. DNA is quantified by PICOGREEN reagent (Molecular Probes) as described above. Samples with low DNA recoveries are reamplified using the same conditions as described above. Samples are diluted with 20% dimethysulfoxide (1:2, v/v), and sequenced using DYENAMIC energy transfer sequencing primers and the DYENAMIC DIRECT kit (Amersham Pharmacia Biotech) or the ABI PRISM BIGDYE Terminator cycle sequencing ready reaction kit (PE Biosystems)

In like manner, the sptm is used to obtain regulatory sequences (promoters, introns, and enhancers) using the procedure above, oligonucleotides designed for such extension, and an appropriate genomic library.

35 IX. Labeling f Pr bes and S uthern Hybridization Analyses

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Microarrays are UV-crosslinked using a STRATALINKER UV-crosslinker (Stratagene). Microarrays are washed at room temperature once in 0.2% SDS and three times in distilled water. Non-specific binding sites are blocked by incubation of microarrays in 0.2% casein in phosphate buffered saline (PBS) (Tropix, Inc., Bedford, MA) for 30 minutes at 60°C followed by washes in 0.2% SDS and distilled water as before.

Hybridization

Hybridization reactions contain 9 μ l of probe mixture consisting of 0.2 μ g each of Cy3 and Cy5 labeled cDNA synthesis products in 5X SSC, 0.2% SDS hybridization buffer. The probe mixture is heated to 65°C for 5 minutes and is aliquoted onto the microarray surface and covered with an 1.8 cm² coverslip. The arrays are transferred to a waterproof chamber having a cavity just slightly larger than a microscope slide. The chamber is kept at 100% humidity internally by the addition of 140 μ l of 5x SSC in a corner of the chamber. The chamber containing the arrays is incubated for about 6.5 hours at 60°C. The arrays are washed for 10 min at 45°C in a first wash buffer (1X SSC, 0.1% SDS), three times for 10 minutes each at 45°C in a second wash buffer (0.1X SSC), and dried.

15 Detection

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Reporter-labeled hybridization complexes are detected with a microscope equipped with an Innova 70 mixed gas 10 W laser (Coherent, Inc., Santa Clara CA) capable of generating spectral lines at 488 nm for excitation of Cy3 and at 632 nm for excitation of Cy5. The excitation laser light is focused on the array using a 20X microscope objective (Nikon, Inc., Melville NY). The slide containing the array is placed on a computer-controlled X-Y stage on the microscope and raster-scanned past the objective. The 1.8 cm x 1.8 cm array used in the present example is scanned with a resolution of 20 micrometers.

In two separate scans, a mixed gas multiline laser excites the two fluorophores sequentially. Emitted light is split, based on wavelength, into two photomultiplier tube detectors (PMT R1477, Hamamatsu Photonics Systems, Bridgewater NJ) corresponding to the two fluorophores. Appropriate filters positioned between the array and the photomultiplier tubes are used to filter the signals. The emission maxima of the fluorophores used are 565 nm for Cy3 and 650 nm for Cy5. Each array is typically scanned twice, one scan per fluorophore using the appropriate filters at the laser source, although the apparatus is capable of recording the spectra from both fluorophores simultaneously.

The sensitivity of the scans is typically calibrated using the signal intensity generated by a cDNA control species added to the probe mix at a known concentration. A specific location on the array contains a complementary DNA sequence, allowing the intensity of the signal at that location to be correlated with a weight ratio of hybridizing species of 1:100,000. When two probes from different sources (e.g., representing test and control cells), each labeled with a different fluorophore, are hybridized to a single array for the purpose of identifying genes that are differentially expressed, the

Probe Preparation from Tissue or Cell Samples

Total RNA is isolated from tissue samples using the guanidinium thiocyanate method and polyA+ RNA is purified using the oligo (dT) cellulose method. Each polyA+ RNA sample is reverse transcribed using MMLV reverse-transcriptase, 0.05 pg/ μ l oligo-dT primer (21mer), 1X first strand buffer, 0.03 units/ μ l RNase inhibitor, 500 μ M dATP, 500 μ M dGTP, 500 μ M dTTP, 40 μ M dCTP, 40 μM dCTP-Cy3 (BDS) or dCTP-Cy5 (Amersham Pharmacia Biotech). The reverse transcription reaction is performed in a 25 ml volume containing 200 ng polyA+ RNA with GEMBRIGHT kits (Incyte). Specific control polyA+ RNAs are synthesized by in vitro transcription from non-coding yeast genomic DNA (W. Lei, unpublished). As quantitative controls, the control mRNAs at 0.002 ng, 0.02 ng, 0.2 ng, and 2 ng are diluted into reverse transcription reaction at ratios of 1:100,000, 1:10,000, 10 1:1000, 1:100 (w/w) to sample mRNA respectively. The control mRNAs are diluted into reverse transcription reaction at ratios of 1:3, 3:1, 1:10, 10:1, 1:25, 25:1 (w/w) to sample mRNA differential expression patterns. After incubation at 37°C for 2 hr, each reaction sample (one with Cy3 and another with Cy5 labeling) is treated with 2.5 ml of 0.5M sodium hydroxide and incubated for 20 minutes at 85°C to the stop the reaction and degrade the RNA. Probes are purified using two successive CHROMA SPIN 30 gel filtration spin columns (CLONTECH Laboratories, Inc. (CLONTECH), Palo Alto CA) and after combining, both reaction samples are ethanol precipitated using 1 ml of glycogen (1 mg/ml), 60 ml sodium acetate, and 300 ml of 100% ethanol. The probe is then dried to completion using a SpeedVAC (Savant Instruments Inc., Holbrook NY) and resuspended in 14 μ l 5X SSC/0.2% 20 SDS. Microarray Preparation

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Sequences of the present invention are used to generate array elements. Each array element is amplified from bacterial cells containing vectors with cloned cDNA inserts. PCR amplification uses primers complementary to the vector sequences flanking the cDNA insert. Array elements are amplified in thirty cycles of PCR from an initial quantity of 1-2 ng to a final quantity greater than 5 μ g. Amplified array elements are then purified using SEPHACRYL-400 (Amersham Pharmacia Biotech).

Purified array elements are immobilized on polymer-coated glass slides. Glass microscope slides (Corning) are cleaned by ultrasound in 0.1% SDS and acetone, with extensive distilled water washes between and after treatments. Glass slides are etched in 4% hydrofluoric acid (VWR Scientific Products Corporation (VWR), West Chester, PA), washed extensively in distilled water, and coated with 0.05% aminopropyl silane (Sigma) in 95% ethanol. Coated slides are cured in a 110°C oven.

Array elements are applied to the coated glass substrate using a procedure described in US Patent No. 5,807,522, incorporated herein by reference. 1 μ l of the array element DNA, at an average concentration of 100 ng/ μ l, is loaded into the open capillary printing element by a high-speed robotic apparatus. The apparatus then deposits about 5 nl of array element sample per slide.

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transposition involving transfer plasmid intermediates. Viral infectivity is maintained and the strong polyhedrin promoter drives high levels of cDNA transcription. Recombinant baculovirus is used to infect Spodoptera frugiperda (Sf9) insect cells in most cases, or human hepatocytes, in some cases. Infection of the latter requires additional genetic modifications to baculovirus. (See e.g., Engelhard,

In most expression systems, SPTM is synthesized as a fusion protein with, e.g., glutathione Ssupra; and Sandig, supra.) transferase (GST) or a peptide epitope tag, such as FLAG or 6-His, permitting rapid, single-step, affinity-based purification of recombinant fusion protein from crude cell lysates. GST, a 26-kilodalton enzyme from Schistosoma japonicum, enables the purification of fusion proteins on immobilized glutathione under conditions that maintain protein activity and antigenicity (Amersham Pharmacia Biotech). Following purification, the GST moiety can be proteolytically cleaved from SPTM at specifically engineered sites. FLAG, an 8-amino acid peptide, enables immunoaffinity purification using commercially available monoclonal and polyclonal anti-FLAG antibodies (Eastman Kodak Company, Rochester NY). 6-His, a stretch of six consecutive histidine residues, enables purification on metal-chelate resins (QIAGEN). Methods for protein expression and purification are discussed in Ausubel (1995, supra, Chapters 10 and 16). Purified SPTM obtained by these methods can be used directly in the following activity assay.

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XIV. Demonstration of SPTM Activity An assay for SPTM activity measures the expression of SPTM on the cell surface. cDNA encoding SPTM is subcloned into an appropriate mammalian expression vector suitable for high levels of cDNA expression. The resulting construct is transfected into a nonhuman cell line such as NIH3T3. Cell surface proteins are labeled with biotin using methods known in the art. Immunoprecipitations are performed using SPTM-specific antibodies, and immunoprecipitated samples are analyzed using SDS-PAGE and immunoblotting techniques. The ratio of labeled immunoprecipitant to unlabeled immunoprecipitant is proportional to the amount of SPTM expressed on the cell surface.

Alternatively, an assay for SPTM activity measures the amount of SPTM in secretory, membrane-bound organelles. Transfected cells as described above are harvested and lysed. The lysate is fractionated using methods known to those of skill in the art, for example, sucrose gradient ultracentrifugation. Such methods allow the isolation of subcellular components such as the Golgi apparatus, ER, small membrane-bound vesicles, and other secretory organelles. Immunoprecipitations from fractionated and total cell lysates are performed using SPTM-specific antibodies, and immunoprecipitated samples are analyzed using SDS-PAGE and immunoblotting techniques. The concentration of SPTM in secretory organelles relative to SPTM in total cell lysate is proportional to the amount of SPTM in transit through the secretory pathway.

Functi nal Assays XV.

calibration is done by labeling samples of the calibrating cDNA with the two fluorophores and adding identical amounts of each to the hybridization mixture.

The output of the photomultiplier tube is digitized using a 12-bit RTI-835H analog-to-digital (A/D) conversion board (Analog Devices, Inc., Norwood, MA) installed in an IBM-compatible PC computer. The digitized data are displayed as an image where the signal intensity is mapped using a linear 20-color transformation to a pseudocolor scale ranging from blue (low signal) to red (high signal). The data is also analyzed quantitatively. Where two different fluorophores are excited and measured simultaneously, the data are first corrected for optical crosstalk (due to overlapping emission spectra) between the fluorophores using each fluorophore's emission spectrum.

A grid is superimposed over the fluorescence signal image such that the signal from each spot is centered in each element of the grid. The fluorescence signal within each element is then integrated to obtain a numerical value corresponding to the average intensity of the signal. The software used for signal analysis is the GEMTOOLS gene expression analysis program (Incyte).

Complementary Nucleic Acids

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Sequences complementary to the sptm are used to detect, decrease, or inhibit expression of the 15 naturally occurring nucleotide. The use of oligonucleotides comprising from about 15 to 30 base pairs is typical in the art. However, smaller or larger sequence fragments can also be used. Appropriate oligonucleotides are designed from the sptm using OLIGO 4.06 software (National Biosciences) or other appropriate programs and are synthesized using methods standard in the art or ordered from a commercial supplier. To inhibit transcription, a complementary oligonucleotide is designed from the most unique 5' sequence and used to prevent transcription factor binding to the promoter sequence. To inhibit translation, a complementary oligonucleotide is designed to prevent ribosomal binding and processing of the transcript. XIII. Expression of SPTM

25 Expression and purification of SPTM is accomplished using bacterial or virus-based expression systems. For expression of SPTM in bacteria, cDNA is subcloned into an appropriate vector containing an antibiotic resistance gene and an inducible promoter that directs high levels of cDNA transcription. Examples of such promoters include, but are not limited to, the trp-lac (tac) hybrid promoter and the T5 or T7 bacteriophage promoter in conjunction with the lac operator regulatory element. Recombinant vectors are transformed into suitable bacterial hosts, e.g., BL21(DE3). Antibiotic resistant bacteria express SPTM upon induction with isopropyl beta-Dthiogalactopyranoside (IPTG). Expression of SPTM in eukaryotic cells is achieved by infecting insect or mammalian cell lines with recombinant Autographica californica nuclear polyhedrosis virus (AcMNPV), commonly known as baculovirus. The nonessential polyhedrin gene of baculovirus is replaced with cDNA encoding SPTM by either homologous recombination or bacterial-mediated 35



Alternatively, the SPTM amino acid sequence is analyzed using LASERGENE software (DNASTAR) to determine regions of high immunogenicity, and a corresponding peptide is synthesized and used to raise antibodies by means known to those of skill in the art. Methods for selection of appropriate epitopes, such as those near the C-terminus or in hydrophilic regions are well described in the art. (See, e.g., Ausubel, 1995, supra, Chapter 11.)

Typically, peptides 15 residues in length are synthesized using an ABI 431A peptide synthesizer (PE Biosystems) using fmoc-chemistry and coupled to KLH (Sigma) by reaction with Nmaleimidobenzoyl-N-hydroxysuccinimide ester (MBS) to increase immunogenicity. (See, e.g., Ausubel, supra.) Rabbits are immunized with the peptide-KLH complex in complete Freund's adjuvant. Resulting antisera are tested for antipeptide activity by, for example, binding the peptide to plastic, blocking with 1% BSA, reacting with rabbit antisera, washing, and reacting with radioiodinated goat anti-rabbit IgG. Antisera with antipeptide activity are tested for anti-SPTM activity using protocols well known in the art, including ELISA, RIA, and immunoblotting.

XVII. Purification of Naturally Occurring SPTM Using Specific Antibodies

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Naturally occurring or recombinant SPTM is substantially purified by immunoaffinity chromatography using antibodies specific for SPTM. An immunoaffinity column is constructed by covalently coupling anti-SPTM antibody to an activated chromatographic resin, such as CNBr-activated SEPHAROSE (Amersham Pharmacia Biotech). After the coupling, the resin is blocked and washed according to the manufacturer's instructions.

Media containing SPTM are passed over the immunoaffinity column, and the column is washed under conditions that allow the preferential absorbance of SPTM (e.g., high ionic strength buffers in the presence of detergent). The column is eluted under conditions that disrupt antibody/SPTM binding (e.g., a buffer of pH 2 to pH 3, or a high concentration of a chaotrope, such as urea or thiocyanate ion), and SPTM is collected.

XVIII. Identification of Molecules Which Interact with SPTM

SPTM, or biologically active fragments thereof, are labeled with ¹²⁵I Bolton-Hunter reagent. (See, e.g., Bolton, A.E. and W.M. Hunter (1973) Biochem. J. 133:529-539.) Candidate molecules previously arrayed in the wells of a multi-well plate are incubated with the labeled SPTM, washed, and any wells with labeled SPTM complex are assayed. Data obtained using different concentrations of SPTM are used to calculate values for the number, affinity, and association of SPTM with the candidate molecules.

Alternatively, molecules interacting with SPTM are analyzed using the yeast two-hybrid system as described in Fields, S. and O. Song (1989) Nature 340:245-246, or using commercially available kits based on the two-hybrid system, such as the MATCHMAKER system (CLONTECH).

SPTM function is assessed by expressing sptm at physiologically elevated levels in mammalian cell culture systems. cDNA is subcloned into a mammalian expression vector containing a strong promoter that drives high levels of cDNA expression. Vectors of choice include pCMV SPORT (Life Technologies) and pCR3.1 (Invitrogen Corporation, Carlsbad CA), both of which contain the cytomegalovirus promoter. 5-10 µg of recombinant vector are transiently transfected into a human cell line, preferably of endothelial or hematopoietic origin, using either liposome formulations or electroporation. 1-2 µg of an additional plasmid containing sequences encoding a marker protein are co-transfected.

Expression of a marker protein provides a means to distinguish transfected cells from

nontransfected cells and is a reliable predictor of cDNA expression from the recombinant vector.

Marker proteins of choice include, e.g., Green Fluorescent Protein (GFP; CLONTECH), CD64, or a

CD64-GFP fusion protein. Flow cytometry (FCM), an automated laser optics-based technique, is used to identify transfected cells expressing GFP or CD64-GFP and to evaluate the apoptotic state of the cells and other cellular properties.

FCM detects and quantifies the uptake of fluorescent molecules that diagnose events preceding or coincident with cell death. These events include changes in nuclear DNA content as measured by staining of DNA with propidium iodide; changes in cell size and granularity as measured by forward light scatter and 90 degree side light scatter; down-regulation of DNA synthesis as measured by decrease in bromodeoxyuridine uptake; alterations in expression of cell surface and intracellular proteins as measured by reactivity with specific antibodies; and alterations in plasma membrane composition as measured by the binding of fluorescein-conjugated Annexin V protein to the cell surface. Methods in flow cytometry are discussed in Ormerod, M. G. (1994) Flow Cytometry, Oxford, New York NY.

The influence of SPTM on gene expression can be assessed using highly purified populations of cells transfected with sequences encoding SPTM and either CD64 or CD64-GFP. CD64 and CD64-GFP are expressed on the surface of transfected cells and bind to conserved regions of human immunoglobulin G (IgG). Transfected cells are efficiently separated from nontransfected cells using magnetic beads coated with either human IgG or antibody against CD64 (DYNAL, Inc., Lake Success NY). mRNA can be purified from the cells using methods well known by those of skill in the art.

Expression of mRNA encoding SPTM and other genes of interest can be analyzed by northern analysis

XVI. Production of Antibodies

SPTM substantially purified using polyacrylamide gel electrophoresis (PAGE; see, e.g., Harrington, M.G. (1990) Methods Enzymol. 182:488-495), or other purification techniques, is used to immunize rabbits and to produce antibodies using standard protocols.

Table 1

				_	_	1 able 1			_	_			
	SE	Q Template II	O Star	t Stop	Frame	Domain	SEQ	Template I	\mathbf{D}	Start	Stop	Fram	Domain
	ID	NO				Type	ID N	O			_		Type
	1	198450.6.oct	272	343	forward 2	TM	26	231583.3.de	вс	1159	1239	forward 1	TM
	1	198450.6.oct	269	334	forward 2	SP		231583.3.de			1233	forward 1	SP
	1	198450.6.oct			forward 2	TM		231583.3.de			1238	forward 3	TM
	2	475178.1.oct	-	1292	forward 3	SP		231583.3.de				forward 1	TM
	2	475178.1.oct	95	172	forward 2	SP		231583.3.de			1233		
				1274								forward 1	TM
		475178.1.oct			forward 3	SP		231583.3.de			1227	forward 1	SP
	2	475178.1.oct	95	157	forward 2	SP		231583.3.de		571		forward 1	SP
	3	231793.2.oct		801	forward 1	SP		231583.3.de		1195	1251	forward 1	TM
-	3	231793.2.oct	739	810	forward 1	SP	26	231583.3.de	₽Ç	1184	1243	forward 2	TM
	3	231793.2.oct	865	930	forward 1	SP	. 26	231583.3.de	ec	1170	1232	forward 3	TM
	3	231793.2.oct	739	810	forward 1	SP	26	231583.3.de	ЭС	1182	1238	forward 3	TM
	3	231793.2.oct	730	810	forward 1	SP.	27	215051.5.de		975	_	forward 3	TM
	4	000010.4.oct		1684	forward 2	SP		215051.5.de			1487	forward 3	TM
	4	000010.4.oct		1696	forward 2	SP		215051.5.de			1492		SP
	5	412959.6.oct		409	forward 2	TM		215051.5.de			1034		SP
	5	412959.6.oct	586	642	forward 1	SP							
	5			406	_			215051.5.de		1394	_	forward 2	TM
		412959.6.oct	350		forward 2	TM		215051.5.de		1424		forward 2	SP
	6	331521.5.oct	807	860	forward 3	TM		215051.5.de			920	forward 3	TM
		331521.5.oct		902	forward 3	SP		215051.5.de		51	140	forward 3	SP
	7	902114.1.oct	288	341	forward 3	SP	27	215051.5.de	€C	506	577	forward 2	SP
	7	902114.1.oct		338	forward 3	SP	27	215051.5.de	ec	1421	1501	forward 2	TM
•	7	902114.1.oct	288	353	forward 3	SP	27	215051.5.de	ec	1424	1480	forward 2	TM
•	7	902114.1.oct	288	347	forward 3	SP	27	215051.5.de	эс	1412	1462	forward 2	TM
- 1	В	481382.1.oct	730	798	forward 1	SP	27	215051.5.de	ec	1424	1471	forward 2	SP
- 1	В	481382.1.oct	730	789	forward 1	SP		215051.5.de			1480	forward 2	SP
		903849.1.oct		1414	forward 2	TM		277726.5.de		= = '	711	forward 1	TM
	9	903849.1.oct		1403	forward 3	SP		277726.5.de			918	forward 1	TM
		433776.4.oct	737		forward 2	SP		277726.5.de			900	forward 1	
		433776.4.oct	797		_	SP						_	TM
		-			forward 2			277726.5.de			426	forward 1	TM
		407607.4.oct		1687	forward 2	TM		277726.5.de		652	729	forward 1	TM
		407607.4.oct		1500	forward 1	SP		277726.5.de			894	forward 1	TM
		234828.6.oct		1180	forward 2	SP		277726.5.de			1430	forward 3	TM
		234828.6.oct		1189	forward 2	SP	28	277726.5.de	ec		903	forward 1	TM
		336430.2.dec		1355	forward 3	SP	28	277726.5.de	BC	844	894	forward 1	TM
	13	336430.2.dec	857	931	forward 2	SP	29	978637.1.de	ec	19	123	forward 1	SP
	13	336430.2.dec	749	850	forward 2	SP	30	240518.12.0	dec	61	114	forward 1	TM
	14	242269.2.dec	769	837	forward 1	TM	30	240518.12.0	dec	64	126	forward 1	TM
	15	432120.2.dec	503	559	forward 2	TM	30	240518.12.0	dec	868	978	forward 1	SP
	16	198060.6.dec	40	126	forward 1	SP	30	240518.12.0	dec		978	forward 1	SP
	17	460295.5.dec	369	449	forward 3	TM	31	413231.8.de	BC.	1182		forward 3	SP
	18	235983.6.dec		3375	forward 1	TM	_	413231.8.de		2531		forward 2	TM
		235983.6.dec			forward 3	SP		413231.8.de			1256	forward 3	SP
		235983.6.dec		3565	forward 2	SP		413231.8.de		-		forward 1	TM
		235983.6.dec			forward 1	TM		413231.8.de					
		235983.6.dec			forward 2	SP						forward 3	SP
		235983.6.dec			forward 2			413231.8.de				forward 3	SP
						TM		413231.8.de				forward 3	SP
		235983.6.dec			forward 2	SP		334406.5.de		886		forward 1	SP
		235983.6.dec		4435	forward 2	SP		411429.8.de		468		forward 3	TM
		235983.6.dec			forward 2	SP	34	320674.7.de	ec	1649	1717	forward 2	TM
	18	235983.6.dec			forward 2	SP	35	197267.1.de	ec	5	76	forward 2	SP
	18	235983.6.dec	4361	4420	forward 2	SP	35	197267.1.de	ЭС	14 .	58	forward 2	SP
	19	238703.2.dec	1057	1140	forward 1	SP	·35	197267.1.de	ec	5	67	forward 2	SP
:	20	038751.5.dec	744	809	forward 3	TM	35	197267.1.de	ec .		67	forward 2	SP
:	20	038751.5.dec	167	238	forward 2	TM		197267.1.de			67	forward 2	SP
		038751.5.dec		803	forward 3	SP		197267.1.de			803	forward 3	SP
		038751.5.dec	464		forward 2	TM		332335.1.de			883	forward 2	SP
		236099.4.dec		1352	forward 3	SP		238992.13.0					
		350875.2.dec	479		forward 2						994	forward 2	SP
						TM		199736.1.de			219	forward 1	TM
		466521.5.dec		666	forward 1	SP		199736.1.de	-		204	forward 1	TM
		466521.6.dec		787	forward 2	SP		199736.1.d			228	forward 1	TM
		474522.8.d c		566	forward 3	SP		228864.5.de			642	forward 1	SP
		474522.8.dec		557	forward 3	SP		228864.5.de		26	139	forward 2	SP
		474522.8.dec		566	forward 3	SP		986539.1.de		3	95	forward 3	SP
		474522.8.dec	507		forward 3	SP ·		481454.4.de		561	647	forward 3	SP
-	26	231583.3.dec	1186	1230	forward 1	TM	41	481454.4.d	С	1239	1298	forward 3	SP

SPTM may also be used in the PATHCALLING process (CuraGen Corp., New Haven CT) which employs the yeast two-hybrid system in a high-throughput manner to determine all interactions between the proteins encoded by two large libraries of genes (Nandabalan, K. et al. (2000) U.S. Patent No. 6,057,101).

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All publications and patents mentioned in the above specification are herein incorporated by reference. Various modifications and variations of the described method and system of the invention will be apparent to those skilled in the art without departing from the scope and spirit of the invention. Although the invention has been described in connection with specific preferred embodiments, it should be understood that the invention as claimed should not be unduly limited to such specific embodiments. Indeed, various modifications of the above-described modes for carrying out the invention which are obvious to those skilled in the field of molecular biology or related fields are intended to be within the scope of the following claims.



SEQ					SEQ				
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1	198450.6.oct	3765347H1	1	286	1	198450.6.oct	2730946H1	1352	1602
1	198450.6.oct	2881536H1	23	287	1	198450.6.oct	g2111659	1365	1747
1	198450.6.oct	2881536F6	23	509	1	198450.6.oct	4501340H1	1369	1614
1	198450.6.oct	3692305H1	226	529	1	198450.6.oct	g3755006	1372	1738
1	198450.6.oct 198450.6.oct	4212539H1	301 343	546 506	1	198450.6.oct	2364954H1	1388	1447
i	198450.6.oct	3451630H1 2614961F6	401	596 999	. 1	198450.6.oct 198450.6.oct	2364915H1 2364954F6	1388 1388	1447 1447
i	198450.6.oct	2614961H1	401	667	1	198450.6.oct	g1238074	1436	1739
1	198450.6.oct	g2111714	487	878	i	198450.6.oct	g2555317	1462	1743
1	198450.6.oct	g1984142	491	674	1	198450.6.oct	g1489886	1468	1745
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1	198450.6.oct	3109843H1	496	663	1	198450.6.oct	3665959H1	1471	1702
1	198450.6.oct	3109585H1	496	595	1	198450.6.oct	4872588H1	1482	1738
1	198450.6.oct	1897893H1	551	795	1	198450.6.oct	g2435210	1490	1739
1 ·	198450.6.oct	1897530H1	551	776	1	198450.6.oct	3133936F6	1516	1734
1	198450.6.oct	5136026H1	599	872	1	198450.6.oct	3133936H1	1517	1755
1	198450.6.oct 198450.6.oct	4505381H1 5894055H1	627 632	891	1	198450.6.oct	1293778H1	1523	1739
1	198450.6.oct	5901947H1	632	895 939	1	198450.6.oct 198450.6.oct	g1497107 3691633H1	1527 1538	1739 1730
i	198450.6.oct	5698273H1	638	908	i	198450.6.oct	q2255347	1554	1809
i	198450.6.oct	3139196H1	638	911	i	198450.6.oct	q3899575	1589	1742
1	198450.6.oct	g1278047	644	1069	i	198450.6.oct	g3735471	1615	1744
1	198450.6.oct	3945278H1	700	978	2	475178.1.oct	g3109369	893	1334
1	198450.6.oct	3941342H1	700	984	2	475178.1.oct	g2106854	903	1285
1	198450.6.oct	g2015083	709	1013	2	475178.1.oct	g1886488	906	1336
1	198450.6.oct	5467754H1	745	1017	2	475178.1.oct	g4243834	921	1335
1	198450.6.oct	g1423015	756	1149	2	475178.1.oct	g3229982	932	1335
1	198450.6.oct	g1497157	778	1271	2	475178.1.oct	g2876106	933	1289
1	198450.6.oct 198450.6.oct	3706512H1 4768061H1	781 786	1071 1024	2	475178.1.oct	g2675531	965	1330
1	198450.6.oct	3683925H1	795	1024	2	475178.1.oct 475178.1.oct	g1479394 g2106969	1023 1028	1335 1335
i	198450.6.oct	5188646H1	822	1148	2	475178.1.oct	g2910156	1042	1311
1	198450.6.oct	1456401H1	868	1143	2	475178.1.oct	6430617H1	1054	1226
1	198450.6.oct	3822085H1	887	1153	2	475178.1.oct	6131966H1	1	187
1	198450.6.oct	3451545H1	888	1141	2	475178.1.oct	4341985H1	1	306
1	198450.6.oct	2804083F6	896	1385	2	475178.1.oct	5696250H1	6	102
1	198450.6.oct	2804083H1	896	1154	2	475178.1.oct	g1577126	219	640
1	198450.6.oct	3519856H1	897	1223	2	475178.1.oct	5475396H1	224	472
1	198450.6.oct	2360526H1	911	1160	2	475178.1.oct	493554H1	246	491
1	198450.6.oct	g1489982	939	1229	2	475178.1.oct	493554R6	246	612
1	198450.6.oct 198450.6.oct	2585474H1 1385485H1	986 1017	1236	2	475178.1.oct	265020H1	249	573
i	198450.6.oct	464838H1	1017	1249 1254	2	475178.1.oct 475178.1.oct	5871350H1 g1886599	259 381	554 851
i	198450.6.oct	g4186989	1051	1479	2	475178.1.oct	5613813H1	383	475
1	198450.6.oct	g3932153	1055	1461	2	475178.1.oct	2518964H1	397	655
1	198450.6.oct	2519149H1	1055	1333	2	475178.1.oct	2518964F6	397	858
1	198450.6.oct	3326033H1	1076	1339	2	475178.1.oct	1533513H1	608	802
1	198450.6.oct	4383511H1	1114	1369	2	475178.1.oct	1533513F6	608	1058
1	198450.6.oct	6323272H1	1119	1395	2	475178.1.oct	g3742669	839	1335
1	198450.6.oct	2804083T6	1124	1676	2	475178.1.oct	g1337822	870	1335
1	198450.6.oct	g1367880	1160	1609	2	475178.1.oct	g3280761	876	1335
1	198450.6.oct	g4223520	1164	1461	2	475178.1.oct	g1337821	877	1347
1	198450.6.oct	2859754T6	1186	1698	2	475178.1.oct	g2883442	883	1311
1	198450.6.oct 198450.6.oct	758599H1 2614961T6	1223 1274	1459	2	475178.1.oct	g2876528	895	1339
i	198450.6.oct	g3755770	1280	1702 1741	2 2	475178.1.oct 475178.1.oct	g1474211 g3001396	1181 1216	1336 1278
i	198450.6.oct	g4390433	1284	1740	2	475178.1.oct	g2768096	1218	1278
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1	198450.6.oct	g3721415	1300	1740	3	231793.2.oct	g3765637	2168	2622
1	198450.6.oct	5491460H1	1301	1434	3	231793.2.oct	5094942H1	2167	2412
1	198450.6.oct	6074015H1	1325	1579	3	231793.2.oct	3673817H1	1003	1285
1	198450.6.oct	g1443523	1326	1739	3	231793.2.oct	4580644H1	1173	1318
1	198450.6.oct	g3367015	1332	1745	3	231793.2.oct	g3049752	2237	2604

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41 481454.4.dec 1206 1298 forward 3 SP	Table 1 cont.
41 4814E4 4 J = 101Waiu 3 SP	50 4000T -
AT ARTACA A CONTAIN Z SP	
41 ARIAEA A OIWAIUZ SP	60 000 13.dec 14/9 1538 forward 2
41 481454 4 de 446 505 forward 2 SP	60 05000.0.dec 1080 1127 forward 2
TOTAL AND END I	00 330399.5 dec 1607 4750 /
	00 350399.5 dec 3743 3904 .
	60 350399.5.dec 1856 1010
42 474800.7.dec 337 430 (1944) 2 SP	60 350300 E 101Walu 2
43 427883.13 dec 36 po (orwald) SP	60 350300 E
AA NIONAFA	60 000000 2169 2234 forward 3
45 353271 0 de - 101 101 111 2 1 M	60 05003.3.UEC 2183 2239 forward 2
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40 221686.2 dec 729 700	60 350300 E
4/ 233347.7.dec 972 1046 (1940 2 SP	60 350300 E da 101Wald 1
47 233347.7 dec 397 470 10 Wald 3 SP	61 085712 0 O Walu 2
47 233247 7 de 101 Ward 3 SP	2469 2528 forward 3
47 222277 7 J	c4 00== 1/92 1848 forward 1
47 232247 7 4	01 005/13.2.dec 2481 2540 forward 3
	01 005/13.2 dec 2447 0500 .
	01 005/13.2.dec 2456 2507 /
	61 085713.2 dec 2456 0540
47 233347.7 dec 264 247 10 Wald 2 SP	61 085713.2 dec 2344 2400 10 Walti 2
47 233347.7.dec 264 220 (Wald 3 IM	61 085712 0 O Walu I
47 233347.7.dec 272 220 (Maiu 3 IM	61 085712 0 d
47 233347 7 da	61 005-10-2. USC 125 1/5 forward 2
47 222247 7 J	et 00== 10.2.dec 2456 2512 forward 2
47 232247 7 4	01 005/13.2.dec 2450 2544 /
10 00007.7.dec 264 335 forward 3 cp	01 005/13.2.dec 2540 2500 4
	02 243014.1.dec 701 ecc .
40 230631.3 dec 534 577 / W	62 245014.1 dec 770 900 (Olward 2
40 230631.3.dec 524 574	62 24E014 4 -1 TOTWAIU 2
48 230631.3 dec 1675 1704	62 245014 4 day ==== 000 lolwald 2
49 335146.1 dec 218 074	63 117464 7 dec 785 856 forward 2
49 335176 4 d O'Walu Z IM	60 447 Get 1411 1458 forward 1 c
49 335146 1 de 203 268 forward 2 TM	60 1473 forward 1
50 337160 1 dec 218 274 forward 2 TM	03 11/464.7.dec 1408 1470 forward 1
	03 11/464.7.dec 1031 1000 / 1031
	63 117464.7 dec 231 270 (Maiu) S
01 040341,12 dec 2500 2654 ,	63 117464.7 dec 574 coo lorratu 3 S
31 340341.12.dec 1208 1204 4	63 117464 7 July 10 Wald S
31 340341.12 dec 2580 2660 /	63 117/64 7 J
31 340341.12 dec 2580 2640	63 117464 7 J
	60 447 dec 1939 1992 forward 1 el
	29/5 3040 forward 2
	60 117 Jec 2/40 2808 forward 1 Cr
	03 11/404./.dec 2066 2050 / 01
	03 11/404./.dec 1006 1000 ;
50 40341.12.dec 3712 3768 forward 1 Tax	63 117464.7 dec 1019 1000 101Wald 1 1N
02 420/45.2 dec 110 404 , """	63 117464.7 dec 1020 1000 101Wald SP
53 444839 17 dec 265 240	
54 245000.6 dec 707 ecc	
54 245000.6 dec 806 860 101Wald 2 1M	63 117464 7
54 245000.6.dec and are 101 Walt 2 SP	302 forward 2 CD
54 245000 6 d - 10 Walu 2 SP	30 1423 1494 forward 1
54 9/Enno o d	00 17704.7.08C 1933 1983 forward 1 The
54 2 5000.0.dec 563 619 forward 2 50	
5/2 634 forward 2 TA	1933 1995 forward 1 TM
	• •
54 445000.6.dec 767 044 / 101	
54 245000.6.dec 812 eec ,	•
57 245000.6 dec 806 eso , 1111	
54 245000.6 dec 772 acc	
54 245000 c d lolwaid 2 Sp	•
56 480710 10 de 270 326 forward 3 TM	
30 400/10.12.dec 2216 2000 /	
5/ 234137.10 dec 549 640	•
30 400530 4 dec 901 001	
39 460951.5.d c 964 1022 t	*
1023 forward 1 SP	
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Table 2 cont. 4 000010.4.oct 4984706H1 1140 1346 000010.4.oct 3165805H1 920 1199 4 000010.4.oct 4594569H1 1142 1396 000010.4.oct 3165853H1 920 1196 4 000010.4.oct 3045057H1 1147 1434 4 000010.4.oct 4729385H1 924 1181 4 4 1401088H1 1166 1403 000010.4.oct 000010.4.oct 6298177H1 937 1259 4 4 000010.4.oct 3552750H1 1172 1391 000010.4.oct 4398053H1 989 1244 4 000010.4.oct 912266H1 1180 1310 4 000010.4.oct 4399305H1 988 1259 4 4 000010.4.oct 1850668T6 1179 1772 000010.4.oct 3600193H1 991 1273 4 000010.4.oct 1326577H1 1209 1470 4 000010.4.oct 4218371H1 991 1165 4 000010.4.oct 000010.4.oct 1323667H1 1209 1430 4 2533935F6 1 130 4 000010.4.oct 984923R1 1211 1664 4 000010.4.oct 2533935H1 212 1 4 000010.4.oct 984923H1 4 1211 1418 000010.4.oct g2205347 30 247 4 000010.4.oct 1261381T6 1223 1769 4 000010.4.oct 2497994F6 30 424 4 000010.4.oct 3818091H1 1225 1478 4 000010.4.oct 5870230H1 28 141 4 000010.4.oct 3819406H1 1225 1495 4 000010.4.oct g2205289 30 267 4 000010.4.oct 3415179H1 1228 1481 4 000010.4.oct 2497994H1 30 145 4 000010.4.oct g1489942 1228 1599 4 000010.4.oct g2002304 30 306 4 000010.4.oct 1234 4357827H1 1489 4 000010.4.oct 149399H1 42 206 4 000010.4.oct 3816557H1 1236 1530 4 000010.4.oct 4421431H1 45 243 4 000010.4.oct g1549922 4 1252 1602 000010.4.oct 5086835H1 59 109 4 000010.4.oct 1960790T6 1276 1765 4 000010.4.oct 3798389H1 74 336 4 000010.4.oct 5621050H1 1288 1573 4 000010.4.oct 2502489F6 79 482 4 000010.4.oct 2502489T6 1294 1765 4 000010.4.oct 2502489H1 294 79 4 000010.4.oct 4297766H1 4 1294 1506 000010.4.oct 3347907H1 95 324 4 000010.4.oct 4298064H1 1294 1534 4 000010.4.oct 1418374H1 95 240 4 000010.4.oct 1843689H1 1305 1561 4 000010.4.oct g2002139 99 536 4 000010.4.oct q518355 1307 1808 4 000010.4.oct 2096696H1 106 351 4 000010.4.oct 2557004H2 274 386 4 000010.4.oct 4911182H1 109 403 4 000010.4.oct 2557066H1 274 518 4 000010.4.oct 1003086H1 343 114 4 000010.4.oct 1816539H1 275 524 4 000010.4.oct 3321596H2 128 373 4 000010.4.oct 3758495H1 283 493 4 3878602H1 129 000010.4.oct 434 4 000010.4.oct 1960790R6 290 364 4 000010.4.oct 3740635H1 129 320 4 000010.4.oct 1960790H1 290 509 4 000010.4.oct 4042307H1 105 163 4 000010.4.oct 2703769H1 314 572 4 000010.4.oct 2587619H1 379 131 4 000010.4.oct 2193493H1 316 571 4 000010.4.oct 2584021H1 131 377 4 000010.4.oct 2878946H1 350 628 4 000010.4.oct 2080578H1 133 385 4 000010.4.oct 4 3674909H1 369 637 000010.4.oct g2410855 145 374 4 000010.4.oct 3957836H2 370 628 4 000010.4.oct 4832261H1 154 326 4 3674529H1 000010.4.oct 370 4 000010.4.oct 613 2226661H1 158 368 4 000010.4.oct 5396777H1 4 385 647 000010.4.oct 996378H1 180 477 4 389 000010.4.oct 4729679H1 662 4 000010.4.oct 3599007H1 181 482 4 000010.4.oct 1232381H1 4 412 649 000010.4.oct 4635262H1 204 450 4 000010.4.oct 2359821H1 418 655 4 000010.4.oct -2716023H1 448 214 4 000010.4.oct 5295590H1 446 716 4 000010.4.oct 3472595H1 232 414 4 000010.4.oct 4667072H1 488 720 4 2557058H1 000010.4.oct 274 523 4 000010.4.oct 2919365H1 528 796 4 000010.4.oct 2497994T6 1577 1762 4 000010.4.oct 5378869H1 547 753 4 000010.4.oct 4127359H1 1588 1807 4 4 000010.4.oct 5398383H1 561 796 000010.4.oct g2444595 1594 1809 4 000010.4.oct 3254113H1 589 4 g2324424 859 000010.4.oct 1594 1808 4 000010.4.oct 4761817H1 592 869 4 000010.4.oct g2324565 1594 1808 4 000010.4.oct 1257093F1 4 599 1194 000010.4.oct 4855237H1 1607 1802 4 000010.4.oct 1257093H1 599 834 4 000010.4.oct 4768170H1 1622 1808 4 000010.4.oct 4984849H1 601 874 4 000010.4.oct 4769968H1 1622 1808 4 000010.4.oct 1850668H1 607 913 4 000010.4.oct 1623 g3133564 1808 4 000010.4.oct 1259277H1 619 861 4 000010.4.oct 5152534H1 1640 1899 4 000010.4.oct 488977H1 625 876 4 000010.4.oct 2533935T6 1677 1766 4 000010.4.oct 5173650H1 632 859 4 000010.4.oct 940189H1 1690 1808 4 000010.4.oct 3990710H1 658 961 4 000010.4.oct 2636973H1 1691 1799 4 000010.4.oct 5072810H1 674 925 4 000010.4.oct 2553110H1 1743 1808 4 000010.4.oct 2995522H1 1042 772 4 000010.4.oct 5002788H1 1752 1808 4 000010.4.oct 4418859H1 781 1037 4 000010.4.oct 5106690H1 1756 1808 4 000010.4.oct 4339619H1 857 1096 4 000010.4.oct 3630725H1 1556 1737 4 000010.4.oct 3207980H1 881 4 991 000010.4.oct 2570529H1 1556 1780 4 000010.4.oct 1261381R1 888 1312 4 000010.4.oct g1717016 1562 1808 4 000010.4.oct 1261381H1 888 1134 5 412959.6.oct 5216131H1 393 673 4 000010.4.oct 1261381R6 888 1341 5 412959.6.oct 5282328H2 399 649 4 000010.4.oct 5714423H1 892 1195 5 412959.6.oct 1286604H1 401 654 4 000010.4.oct 447362H1 900 1120 5 412959.6.oct 5075560H1 403 678 000010.4. ct 5265158H1 5 904 1130 412959.6.oct g4306032 416 815

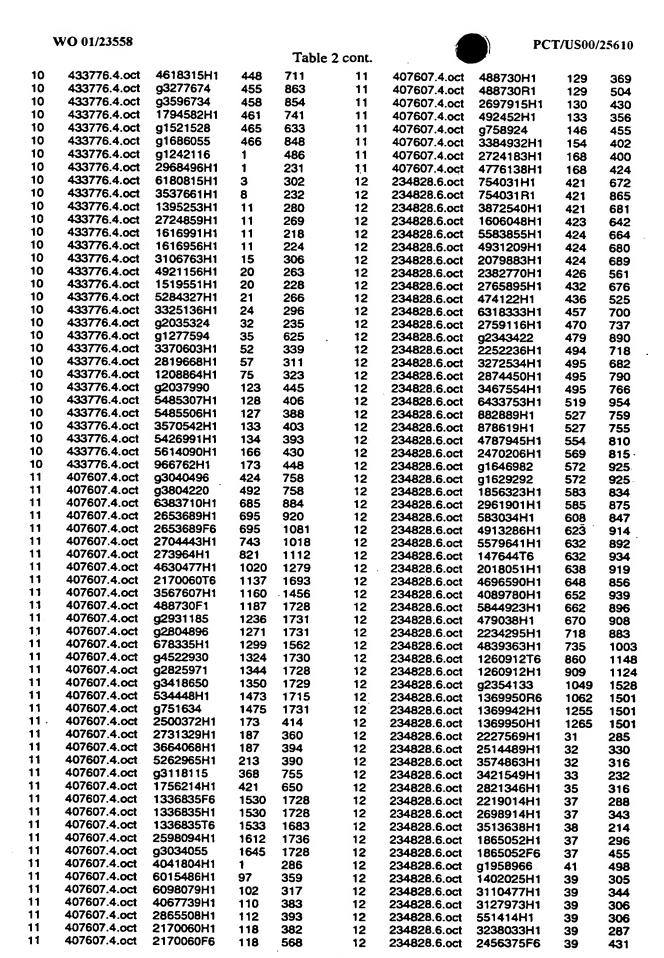
W O 01/23558				•9	
		Table	. 2		PCT/US00/25610
3 231793.2.oct 1	707686T6 22		2 cont.	*	0 17 0500725010
3 231793.2.oct 5			3 23179	3.2.oct 1914969	
3 231793.2 oct 20	27000				H1 2225 2477
3 231793.2 oct of	2000-1-		3 23179		H1 21 200
3 231793.2 oct ~	10700				H1 22 240
	1078219 233		-01100		H2 30 305
	83970T6 236	5 2585			F6 33 444
3 231793.2.oct 37	12992H1 236	5 2563		3.2.oct 1866779	H1 33 340
2 201700.2.001 3/	43582H1 1	299	-01750	3.2.oct 1311083	H1 80 200
2 40.2.001 34	90362H1 3	199		1.2.oct 3236153	H1 111 250
2 1.00.2.001 03	01123H1 10	295		-2.oct 5158979	11 202 425
24	88457H1 117	6 1404		.2.oct 1390306F	11 287 527
20,1,001 20,	23858R6 119		-01730	.2.0Cl 1300242L	14
2 20,4:001 20,	23858H1 1194		400010	.4.oct 32542137	6 1335 1700
2 40	0753H1 1245		4 000010.	4.oct 1370046F	11 1337 1400
231/93.2.oct 279	3766H1 1364		4 000010.	4.001 53067777	4 1400
231/93.2.oct 180	3565H1 1546		4 000010.	4.0Ct 13700460	
231/93.2.oct 617	3678H1 1561		4 000010.	4.oct 1370102W	4
231/93.2.oct 121	4293T6 2183		4 000010.	4.0Ct 1726162E	- 1007
231/93.2.oct 121	4293R6 2184	2577	4 000010.4	1.oct 5372022L	4 40
231793.2.oct 121	4293H1 210A	2407	4 000010.4	oct 1726160U	1073
231/93.2.oct 180	3565T6 2104		4 000010.4	OCT 1726160Te	1049
231/93.2.oct 100	126T6 2002	2547 2558	4 000010.4	Oct 02002504	1770
231/93.2.oct 417(0673H1 2146	2407	4 000010.4	Oct d364E063	1351 1580
231/93.2.oct 5027	277H1 100E		4 000010.4	OCT 2400264To	1355 1808
231/93,2,oct 3803	405H1 2038	2261	4 000010.4	.oct n30510cc	
231/93.2.00 5005	560H1 2046	2335	4 000010.4	Oct 01480042	1379 1808
231/93.2.0ct 4212	438H1 2072	2221	4 000010.4	oct 5085122U4	1390 1808
431/93.2.0ct 1096	603T6 2084	2273	4 000010.4	OCT 04019720	1072
231/93.2.oct 6197	9679 1896	2560	4 000010.4	oct 04060310	1393 1810
3 231793.2.oct 4593	321H1 1918	2192	4 000010.4.	oct 041000e1	1396 1809
23 1/93.2.0ct 5086	230H1 1917	2189	4 000010.4.	oct 03307404	1400 1812
3 231793.2.oct gazar	027 1948	2189	4 000010.4.0	oct 02206554	1406 1817
3 231793,2,oct 6171		2321	4 000010.4.0	oct 1654556H1	1408 1809
3 231793.2.oct 59515	200114	2242	4 000010.4.0		1415 1631
3 231793.2.oct 26730		2290	4 000010.4.0		1416 1580
3 231793,2,oct 26730		2501	4 000010.4.0		1419 1809
3 231793.2.nct 97701	0114	2213	4 000010.4.0		1423 1580
3 231793.2.oct 52815	70110	2200	4 000010.4.0		1429 1672
3 231793.2.oct 47317		1837	4 000010.4.0		1430 1690
3 231793.2.oct 15829		1859	4 000010.4.0		1435 1808
3 231793.2.oct 15828	104	2082	4 000010.4.00		1441 1811
3 231793.2.oct 45089	714		000010.4.00		1456 1636
3 231793.2.oct 355966	CU4 4	1901 4	000010.4.00		1477 1808
3 231793.2.oct 396101	CUI4	1994 4	000010.4.00		1487 1807
3 231793.2 nm 360500	OLIA ATT	835 4	000010.4.00		1495 1807
3 231793.2.oct 404871	0144	042 4	000010.4.00	0-1-UL-4	1496 1809
231/93.2.0ct 170760	01.44	092 4	000010.4.oc		1500 1580
3 231793,2,oct 529600	21.14	072 4	000010.4.oc		1512 1803
3 231793.2.oct 023221	_	102 4	000010.4.oc		1513 1808
3 231793.2.oct 130206	777	604 4	000010.4.oct	956928T1 9821709	1512 1769
3 231793.2.oct 130251	7EC	562 4	000010.4.oct	g2100454	1522 1833
3 231793.2.oct 1302517	PLIA	502 4	000010.4.oct		1521 1808
3 231793.2.oct 1347221	14	502 4	000010.4.oct		1537 1781
3 231793.2.oct 0110571		502 4	000010.4.oct	40.	1544 1808
3 231793.2.oct 1986603	DC -	510 4	000010.4.oct		991 1233
3 231793.2.oct 1991005	LI4 OLD	85 4	000010.4.oct	E40054 4144	1006 1066
3 231793,2,oct 3962341	U4	72 4	000010.4.oct	4000400	1006 1242
3 231793,2.oct 1001126	20 00	58 4	000010.4.oct		1011 1294
3 231793.2.oct 4049712			000010.4.oct		1038 1277
3 231793.2.oct 37671011	14	7 4	000010.4.oct		1058 1361
3 231793.2.oct 27917051	14		000010.4.oct	~1000~~	1064 1333
3 231793.2.oct 45760541	14	7 4	000010.4.oct		1068 1409
3 231793,2,oct 55028201	14 == : 021) 4	000010.4.oct		1069 1257
3 231793.2.oct 5052270	14 330) 4	000010.4.oct		080 1212
3 231793.2. ct 1986602L	1 200	4	000010.4.oct		097 1370
3 231793.2.oct 12141671		4 4	000010.4.oct	1315140H1 1	110 1358
3 231793,2,oct 50847691	1	9 4			110 1402
3 231793.2.oct g2347914		7 4	000010.4.oct	4858515H1 1	122 1353
9234/914	2221 260	1 4		1459590H1 1	124 1376
		<i>C</i> 1	100.4.00	4292624H1 1	134 1395
		61			

				Table 2	cont.				
5	412959.6.oct	g2265153	614	967	6	331521.5.oct	g3736851	1121	1507
5	412959.6.oct	g1745215	619	815	6	331521.5.oct	5598577H1	1123	1315
5	412959.6.oct	g2368862	627	815	6	331521.5.oct	g2270261	1138	1501
5	412959.6.oct	g2703493	628	970	6	331521.5.oct	927617T6	1142	1460
5	412959.6.oct	g1493098	628	982	6	331521.5.oct	g922764	1144	1480
5	412959.6.oct	g3040120	630	978	6	331521.5.oct	g922061	1150	1496
5	412959.6.oct	g2575651	631	815	6	331521.5.oct	g3417656	1150	1498
5	412959.6.oct	g1618594	642	971	6	331521.5.oct	g3745245	1150	1498
5	412959.6.oct	g1148550	649	885	6	331521.5.oct	3943483H1	1035	1301
5	412959.6.oct	2821589H1	656	959	6	331521.5.oct	1787637H1	1041	1273
5	412959.6.oct	g1218727	682 ·	961	6	331521.5.oct	838819H1	634	888
5	412959.6.oct	g787387	692	967	6	331521.5.oct	g1187465	664	913
5	412959.6.oct	1695326H1	691	815	6	331521.5.oct	4667001H1	692	967
5	412959.6.oct	g2177825	702	967	6	331521.5.oct	3113662H1	509	730
5	412959.6.oct	2725050H1	718	815	6	331521.5.oct	g1986323	509	821
5	412959.6.oct	2674048T6	758	933	6	331521.5.oct	2732541H1	516	739
5	412959.6.oct	g1860722	764	1003	6	331521.5.oct	g1239894	558	772
5	412959.6.oct	g1139057	773	982	6	331521.5.oct	3513041H1	557	799
5	412959.6.oct	g3145267	794	949	6	331521.5.oct	3190732H1	607	963
5	412959.6.oct	g1265363	803	975	6	331521.5.oct	2375477H1	505	725
5	412959.6.oct	4771510H1	819	989	6	331521.5.oct	3149260H1	506	788
5	412959.6.oct	778431H1	825	981	6	331521.5.oct	g3595860	1088	1498
5	412959.6.oct	g3149926	832	885	6	331521.5.oct	g3146187	1099	1501
5	412959.6.oct	g3043195	849	970	6	331521.5.oct	4979280H1	1109	1373
6	331521.5.oct	2211724H1	478	739	6	331521.5.oct	g3250147	1114	1505
6	331521.5.oct	3456836H1	480	709	6	331521.5.oct	g3431781	1118	1499
6	331521.5.oct	2202654H1	485	720	6	331521.5.oct	3629028H1	1044	1329
6	331521.5.oct	2661126T6	968	1458	6	331521.5.oct	g3404876	1057	1501
6	331521.5.oct	g3872415	1033	1498	6	331521.5.oct	3444045H1		1315
6	331521.5.oct	3943483F6	1035	1499	6	331521.5.oct	1212854T6	1084	1475
6	331521.5.oct	g3884652	1157	1503	6	331521.5.oct	5994615H1	491	790
6	331521.5.oct	g1197983	1173	1502	6	331521.5.oct	5574629H1	492	750
6	331521.5.oct	4666217T6	896	1475	6	331521.5.oct	3091023H1	490	777
6	331521.5.oct	4378681H1	920	1191	6	331521.5.oct	077179H1	491	690
6	331521.5.oct	5563982H1	951	1165	6	331521.5.oct	422847H1	495	777
6 6	331521.5.oct	4666217F6	891	1066	6	331521.5.oct	g2224124	1204	1502
6	331521.5.oct 331521.5.oct	4666217H1 g3801327	891 1271	1158 1499	6 6	331521.5.oct	g2238189 5730657H1	1216 1233	1516 1488
6	331521.5.oct	g2411273	1285	1501	6	331521.5.oct 331521.5.oct	3510032T7	1233	1459
6	331521.5.oct	g566119	1304	1497	6	331521.5.oct	g2882731	1242	1503
6	331521.5.oct	6092081H1	1329	1498	6	331521.5.oct	g570486	1	181
6	331521.5.oct	853544H1	1352	1485	6	331521.5.oct	g875762	2	309
6	331521.5.oct	2705657T6	1353	1457	6	331521.5.oct	g831148	2	358
6	331521.5.oct	2705657H1	1360	1498	6	331521.5.oct	4175396H1	494	777
6	331521.5.oct	2705657F6	1360	1498	6	331521.5.oct	4123843H1	499	635
6	331521.5.oct	g868758	1416	1505	6	331521.5.oct	781855R1	498	1067
6	331521.5.oct	g907918	1444	1499	6	331521.5.oct	781855H1	498	742
6	331521.5.oct	1212854H1	718	1014	6	331521.5.oct	g1281868	504	951
6	331521.5.oct	5097150H1	718	983	6	331521.5.oct	5463846H1	497	689
6	331521.5.oct	2005993H1	718	889	6	331521.5.oct	5120226H1	506	790
6	331521.5.oct	5114952H1	727	996	6	331521.5.oct	g830914	1182	1505
6	331521.5.oct	4032072H1	780	1019	6	331521.5.oct	g1486742	1197	1517
6	331521.5.oct	2046722H1	445	722	6	331521.5.oct	g4296634	1200	1504
6	331521.5.oct	6436791H1	451	976	7	902114.1.oct	3815354H1	1.	277
6	331521.5.oct	2507802H1	475	713	7	902114.1.oct	6099860H1	3.	269
6	331521.5.oct	3673647H1	431	736	7	902114.1.oct	g3419125	193	601
6	331521.5.oct	g922133	444	781	7	902114.1.oct	1869318H1	205	458
6	331521.5.oct	g922831	441	726	7	902114.1.oct	q3434169	236	599
6	331521.5.oct	3510032F6	255	827	7	902114.1.oct	g2211756	313	614
6	331521.5.oct	3510032H1	255	561	7	902114.1.oct	4664818H1	395	644
6	331521.5.oct	3115530H1	435	730	7	902114.1.oct	934433R1	476	921
6	331521.5.oct	2579471H1	427	706	7	902114.1.oct	934433H1	476	752
6	331521.5.oct	g1156421	709	861	7	902114.1.oct	2848096H1	725	985
6	331521.5.oct	g875763	1152	1505	8	481382.1.oct	4140109H1	35	204
6	331521.5.oct	g2279775	1153	1499	8	481382.1.oct	3793754H1	35	189
6	331521.5.oct	ğ1158127	1154	1503	8	481382.1.oct	2769459H1	35	116
6	331521.5.oct	g4147609	1155	1499	8	481382.1.oct	1732095T6	35	237
6	331521.5.oct	g2240963	1118	1498	8	481382.1.oct	1732095H1	84	358

'		T.I.	D	CT (I I Cook to to
5 412959.6.oct 2470746H	1 416	Table 2 cont.	,	CT/US00/25610
5 412959.6.oct 5353589Ti		14 5 4129 66 5 4100	59.6.oct 2763336H1	
5 44189438	400	5 4129	9.6.oct 170274444	2-73
5 412959.6 ort 34188920		3 4129	9.6.oct 1632693H1	14 000
3 412050 6 604	437 g	41295	9.0.0Ct 4296049H1	14 225 14 268
3 412959.6.oct 03530347	439 8	5 5 41295	9.0.0ct 5074132H1	14 297
	444 81	5 5 41295		1 268
5 412959.6.oct g3048416	445 67 448 81	<u> </u>	9.6.oct 01157175	2 234
5 412959.6.oct g2782788	448 81 447 81	² 3 412959	9.6.oct 125200414	1 358
5 41200.0.001 92901391	464 81	<u> </u>	6.0ct 4527020U+	1 212 1 264
5 412959 6 oct 93801539	471 81	3 412959	.6.oct g2029372	4
5 412959.6.oct gazatora	480 889	412959	.6.oct 775186H1	1 177 2 217
412959.6.oct g2675057	489 979	5 412050		2 480
5 412959.6.oct g3038152	493 815 493 970	5 412959	6.oct 3206555114	1 218
5 412959.6.oct g1639027	493 970 501 715	⁵ 412959	6.0ct 336033414	2 177
5 412950 6 24 943/3288	506 973	o 412959.	6.0ct 3405500114	3 233 14 298
g 1103-0.00t g3988957	509 815	5 412959 5 412959	o.oct 4655833H1	10
5 412959.6.oct 93961978 5 412959.6.oct 93869490	522 815		o.oct 496681H1	16 264 23 259
5 412959,6,oct 04004707	525 968	5 412959. 5 412959.		26 206
5 412959.6.oct g3086363	525 977 530 815	5 412959	6.oct 496697H1	23 248
5 412959.6.oct g3897838	530 815 533 967	o 412959.6	Oct 6110001114	27 292
5 4400-0.001 92115818	534 973	³ 412959.6	OCT 01745450	29 247 29 322
5 412959 6 000 9328 1206	537 970		.oct 3879459H1	29 322 35 312
5 412959.6.oct 04060000	540 971	5 412959.6 5 412959.6	.oct 3752872H1	39 255
5 412959.6.oct gaggagga	543 980	5 412959 6		59 311
5 412959.6.oct g1331787	545 982 545 973	³ 412959.6	OCT 01040400	60 188
5 412959.6.oct g2526614	545 973 546 976	³ 412959.6	Oct 200449014	73 532
5 412959 6 oct 5000	548 974	³ 412959.6.	oct g1987165	130 412 144 469
5 412959.6.oct g2264700	558 805	, ,	oct g1237710	1 44 469 146 287
5 412959.6.oct 03394070	563 967	5 412959.6. 5 412959.6.	oct 2120694H1	166 337
412959.6.oct 2672133H1	573 973 581 814	³ 412959 6 7	ct azecono	165 470
5 412959.6.0ct 1772941H1	581 814 581 855	³ 412959.6.d	ct 981393U4	173 499
g3038159	584 970	3 412959.6.c	Ct 3705420114	191 432 201 468
5 412959 6 oct =007000	591 976	⁵ 412959.6.0	ct g1745306	
412959.6.oct 01741460	597 970	5 412959.6.0 5 412959.6.0	ct g1981799 2	06 502 45 508
5 412959.6.oct g1883800	949	5 412959.6.0	ct 2581258H1 3	39 583
5 4120E0 6 a.s.	967	5 412959.6.o	1 179150770	40 621
5 412959.b.oct 2654578H1 3		⁵ 412959.6.00	t 412004114	
5 4450844H1 A		³ 412959.6 oc	t 500011011	48 631 17 644
5 412959.6.oct 02150400 4	269	5 412959.6.oc 5 412959.6.oc	t 3771103H1 35	~
5 412959.6.oct 406569414		5 412959.6.oc		60 619
5 412959.6.oct 686032H1 7	276	³ 412959 6 cc	202040470	0 826
5 412959.b.oct 3988406H1 7	225 195	³ 412959.6.oct	295610014	
5 412959 6 cm 47056H1 9	275	5 412959.6.oct	01310510	_ •.•
5 412959 6 oct 415 425 11	208		g1447775 25	
5 412959.6.oct 3156241U4	257	5 412959.6.oct 5 412959.6.oct	91882897 250	
5 412959.6.oct 255159614	_00	5 412959 6 oct	23500071	526
5 412959.6.oct g1996909 10	255 311	5 412959.6.oct	120700411	
5 4400-0-0-0-0 U343/5H1 12	226	3 412050 6 004	1507384H1 275 1541305H1 288	
5 412959 6 000 507 13	306	5 412959.6.oct 5 412959.6.oct	4744668H1 392	
5 412959.6.oct 5152592U4	185	5 412959.6.oct 5 412959.6.oct	5435925H1 300	532
5 412959.6.oct 506022214	277	5 412959 6 001	g2838960 866	972
5 412959.6.oct 3706309H1 7	271	5 412959.6.oct	g1745821 866 721875H1 866	967
5 412959.b.oct 1333267H1 14	290 298	5 412959.6.oct	70100011	958
5 412959.6.oct 2764864H1 14	255 255	5 412959.6.oct	93041378 866 93041378	958
5 412959.6.oct 3470540144 14	282		91193527 907	970 979
5 412959.6.oct 2012750114	247	5 412959.6.oct 412959.6.oct	g2185358 599	979 971
5 412959.6.oct 4295435H1 14	286	5 412959.6.oct	93785891 609	995
- 100111 14	234		93015900 610 1918481H1 614	970
	6:		1910481H1 614	815

	•			Table	2 cont.				
9	903849.1.oct	6260688H1	1026	1267	9	903849.1.oct	4245542H1	257	502
9	903849.1.oct	1613848H1	1028	1243	9	903849.1.oct	1450643F1	73	498
9	903849.1.oct	4157804T8	1035	1591	9	903849.1.oct	4245904H1	257	517
9	903849.1.oct	6190485H1	1034	1319	9	903849.1.oct	1450643H1	73	332
9	903849.1.oct	4550411T1	1047	1573	9	903849.1.oct	5698478H1	77	265
9	903849.1.oct	2173189T6	1048	1572	9	903849.1.oct	2539842H1	87	329
9	903849.1.oct	4549428T1	1061	1555	9	903849.1.oct	5065835H1	103	348
9	903849.1.oct	748919H1	1074	1310	9	903849.1.oct	5377044H1	131	394
9	903849.1.oct	748919R1	1074	1611	9	903849.1.oct	3689228H1	292	570
9	903849.1.oct	2700820H1	1083	1352	9	903849.1.oct	g1276068	313	801
9	903849.1.oct	1709866H1	1095	1321	9	903849.1.oct	3479853H1	321	651
9	903849.1.oct	1709866F6	1095	1473	9	903849.1.oct	2951083H1	330	589
9	903849.1.oct	2350024H1	1097	1310	9	903849.1.oct	955261R1	334	807
9	903849.1.oct	2719920H1	1101	1354	9.	903849.1.oct	955261H1	334	615
9	903849.1.oct	2720624H1	1101	1335	9	903849.1.oct	3234424H1	345	517
9	903849.1.oct	3051469H1	1108	1388	9	903849.1.oct	264620H1	358	701
9	903849.1.oct	3050747H1	1108	1446	9	903849.1.oct	2344474F6	369	815
9 9	903849.1.oct	1870888T6	1119	1571	9	903849.1.oct	2344474H1	369	537
9	903849.1.oct 903849.1.oct	1255114T6	1124	1569	9	903849.1.oct	3684753H1	378	693
9		g2575208 g3151314	1135	1619	9	903849.1.oct	3777349H1	390	672
9	903849.1.oct 903849.1.oct	g1018704	1139 1142	1616 1413	9	903849.1.oct 903849.1.oct	2636983H1 3969738H1	398	638
9	903849.1.oct	1709866T6	1148	1579	9	903849.1.oct	3720303H1	403 408	689 700
9	903849.1.oct	g2837554	1150	1614	9	903849.1.oct	4616163H1	414	680
9	903849.1.oct	g1801648	1	220	9	903849.1.oct	1610601H1	417	655
9	903849.1.oct	4768624H1	i	267	9	903849.1.oct	1610601F6	417	797
9	903849.1.oct	3693504H1	i	307	9	903849.1.oct	4325728H1	419	601
9	903849.1.oct	3693519H1	i	301	9	903849.1.oct	3660219H1	422	641
9	903849.1.oct	4160192H1	3	250	9	903849.1.oct	4542103H1	422	675
9	903849.1.oct	3617693H1	3	278	9	903849.1.oct	5210061H1	430	660
9	903849.1.oct	3617093H1	3	254	9	903849.1.oct	1418988H1	431	673
9 ·	903849.1.oct	3651125H1	6	290	9	903849.1.oct	972888H1	431	668
9	903849.1.oct	3381741H1	14	247	9	903849.1.oct	3202649H1	438	664
9	903849.1.oct	4527453H1	16	93	9	903849.1.oct	4637919H1	451	714
9	903849.1.oct	4084909H1	16	192	9	903849.1.oct	3415427H1	464	722
9	903849.1.oct	491926H1	16	197	9	903849.1.oct	4093051H1	477	743
9	903849.1.oct	3792824H1	17	282	9	903849.1.oct	5734663H1	495	746
9	903849.1.oct	3743920H1	18	321	9	903849.1.oct	g944573	510	854
9	903849.1.oct	3460060H1	17	276	9	903849.1.oct	g1694097	515	846
9	903849.1.oct	4527596H1	23	270	9	903849.1.oct	g751975	522	768
9 9	903849.1.oct	3686573H1	19	317	9	903849.1.oct	000134H1	527	983
9	903849.1.oct	2173189F6	20	94	9	903849.1.oct	5207738H1	530	767
9	903849.1.oct 903849.1.oct	2549457H1	18	270	9	903849.1.oct	3092084H1	540	805
9	903849.1.oct	6384945H1 2173189H1	20 20	324 251	9 9	903849.1.oct 903849.1.oct	3092084F6 3877454H1	541 548	982 825
9	903849.1.oct	3510073H1	23	294	9	903849.1.oct	4441451H1	549	794
9	903849.1.oct	3225307H1	23	316	9	903849.1.oct	a920316	589	863
9	903849.1.oct	1870888F6	23	547	9	903849.1.oct	1527791H1	44	242
9	903849.1.oct	1870888H1	23	287	9	903849.1.oct	3074436H1	44	308
9	903849.1.oct	3075151H1	23	317	9	903849.1.oct	g831292	26	423
9	903849.1.oct	2720653H1	23	277	9	903849.1.oct	g1291703	27	510
9	903849.1.oct	3507839H1	24	313	9	903849.1.oct	g573100	36	337
9	903849.1.oct	5842380H1	26	91	9	903849.1.oct	1824463H1	35	263
9	903849.1.oct	1870853H1	25	251	9	903849.1.oct	5117411H1	39	298
. 9	903849.1.oct	4552316H1	24	172	9	903849.1.oct	4984956H1	36	288
9	903849.1.oct	3742953H1	25	324	9	903849.1.oct	483920H1	40	265
9	903849.1.oct	5276481H1	208	377	9	903849.1.oct	4154493H1	41	292
9	903849.1.oct	3284445H1	62	308	9	903849.1.oct	g1274878	43	654
9	903849.1.oct	2108887H1	211	353	9	903849.1.oct	2768516H1	41	302
9	903849.1.oct	2475835H1	63	284	9	903849.1.oct	1460203H1	41	273
9	903849.1.oct	3029734H1	65	343	9	903849.1.oct	g1271819	44	331
9	903849.1.oct	g1959086	65	486	9	903849.1.oct	2259610H1	42	249
9	903849.1.oct	3289141H1	214	445	9	903849.1.oct	3692010H1	46	337
9	903849.1.oct	g1958846	67	406	9	903849.1.oct	4897221H1	45	355
9	903849.1.oct	4246274H1	257	505	9	903849.1.oct	2908015H1	41	266
9	903849.1.oct	2814301H1	72	359	9	903849.1.oct	2403093H1	42	249
9 9	903849.1.oct	485232H1	72	372	9	903849.1.oct	495889H1	43	250
3	903849.1.oct	485973H1	72	31 <u>5</u>		903849.1.oct	3588716H1	42	352

8	491200 4			Ta	ible 2 c	ont.		r	C1/US0	0/25610
8	481382.1.0		129	370						
8	481382.1.0		179	422		9 90384	9.1.oct	2624665H1	1394	400-
8	481382.1.0		206	389		9 90384	9.1.oct (2695471		1608
8	481382.1.0	ct 2508704H1	252			9 90384	9.1.oct 2	2988037	1408	1620
	481382.1.00	Ct 2370122U4	285	489		9 90384	9.1.oct 2	2875731	1413	1610
8	481382.1.oc	237012256	285	508		9 903849	9.1.oct 3	120483H1	1418	1612
. 8	481382.1.oc	# 280303744		610		9 903849		120403H1	1423	1602
8.	481382.1.oc	1 533530550	308	418		9 903849		99503T1	1431	1572
8	481382.1.oc	1 6391567114	325	822		9 903849		99503H1	1431	1610
8	481382.1.oc	1431535H1	334	534		9 903849		680380H2	1439	1589
8	481382.1.oc		378	650		9 903849		1331532	1452	1617
8	481382.1.oc		396	656				2806322	1462	1610
8	481382.1.oct		586	848				3093063	1468	1616
9	903849.1.oct		1	285				3041606	1471	1618
9	903849.1.oct		1180	1621	Š		1.oct gs	20645	1481	
9	903849.1.oct		1191	1421			1.oct 23	25791H1	1511	1617
9	903849.1.00		1195	1610	9		1.oct 23	25782H1		1615
9	903849.1.oct		1200	1567	9		1.0ct aa	087654	4	1608
9	903849.1.oct		1203	1592	9	903849.	1.oct 31	65133H1		1610
9	903849.1.oct	g1803794	1209	1608	9	903849.	1.oct 25	07856H1		874
9	903849.1.oct	1697502H1	1209		9	903849:1	1.0ct a16	570047		878
9	903849.1.oct	g3366973	1210	1417	9	903849.1	oct as	90906		1015
9	903849.1.oct	g3245013	1219	1614	9	903849.1	oct aza	51221		862
9	903849.1.oct	94078415	1223	1611	. 9	903849.1	oct 125	5114F6		365
9	903849.1.oct	g2740706	1225	1614	9	903849.1	oct 125	5114H1		157
	903849.1.oct	92463862		1614	9	903849.1	oct 395	3586H1		397
9	903849.1.oct	g2657445		1612	9	903849.1	oct 291	222211		193
9	903849.1.oct	1970910H1		1610	9	903849.1	oct 618		676 8	155
9	903849.1.oct	g519042	44	1493	9	903849.1		~~~~		46
9	903849.1.oct	g3739697		1610	9	903849.1	oct 20s	~~~		109
9	903849.1.oct	g3306909		1614	9	903849.1.				48
9	903849.1.oct	g2525781		1614	9	903849.1.	oct 1224	COCLA		37
9	903849.1.oct	a12000		1438	9	903849.1.				94
9	903849.1 oct	-5040		1599	9	903849.1.	oct 4703		749 10	024
9	903849.1.oct	1600000000		614	9	903849.1.0			783 10	248
9	903849.1.oct			440	9	903849.1.0		~~~)58 -
9	903849.1.oct	2007400		614,	9	903849.1.0		6808 8		77
9	903849.1.oct	71600040		518	9	903849.1.0		04 - 00	45 92	:6
9	903849.1.oct 2	1100000		582	9	903849.1.0			45 10	76
9 .	903849.1.oct			608	9	903849.1.0		926H1 8	45 9 6	0
9 ;	903849.1.oct	754004	284 1	614	9	903849.1.0		420H1 8	45 10	94
9 9	903849.1.oct	11110000		502	9	903849.1.0			45 13:	
9 9	903849.1.oct	440000		518	9	903849.1.00			31 112	
9 9	03849.1.oct o	1001010		314	9	903849.1.00				32
9 9	103849.1.oct 7	E446444		14	9	903849.1.00		7/H1 87	7 1 101	6
9 9	03849.1.oct a	2470545		98	9	903849.1.00			' ¹ 132	
9 9	03849.1.oct a	1001700		17	9	903849.1.00		66 87	1 109	8
9 9	03849.1.oct G	44040		14	9	903849.1.oc		62H1 87	1 111	0
9 9	03849.1.oct or	1640004	331 16		9	903849.1.oc		86H1 87	3 114	
9 9	03849.1.oct at	1044545	332 16		9	903849.1.oc	18404	1H1 87	6 109	0
9 9	03849.1.oct no	20 45 400	338 16 ⁻		9	903849.1.oct		53H1 878	3 107	
9 90	03849.1.oct 01		39 161		9	903849.1.oct		907 907	7 1163	
9 90		20000-	39 161		9 .	903849.1.oct		6H1 930	127	
9 90	3849.1.oct at	04000.	42 161		9	903849.1.oct		OH1 937	1208	
9 90		4507414		Ю	9	903849.1.oct		H1 939	1165	
9 90		45971H1 13		4	9	903849.1.oct		0H1 940	1226	
9 90		124543 13		0	9	903849.1.00		0F6 940		
9 90		226701 136	61 161			903849.1.oct		H1 948	1205	
9 90:		382614 136	33 161			903849.1.oct		3H1 972	1239	
		3980H1 136	35 1610			903849.1.oct	1929780	T6 977		
		0854 136	55 1605			903849.1.oct	760376F	31 986	1465	
		44590 115	5 1608			903849.1.oct	760376H	11 986	1258	
		7148H1 115				903849.1.oct	g114679	6 1005		
		65122 115				903849.1.oct	9116247	8 1006		
		9110H1 116				903849.1.oct	5598908	H1 1012		
		76029 136				903849.1.oct	3523694	H1 1014		
		17308 136 <i>6</i>			9 9	003849.1.oct	5598808	H1 1014		
	849.1.oct 9750	0565 ₁₃₆₀			9 9	003849.1.oct	4466790	H1 1017		
	849.1.oct g518	3777 ₁₃₇₀			9 9	03849.1.oct	2344474	T6 1010		
903	849.1.oct g136	8048 1379			9 9	03849.1.oct	507835H	1 1022		
			1011	٤	9 9	03849.1.oct	3925430	11 1024	1113	
				65				1024	1224	
				U.J						



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	9 903849	9.1.oct	4558109H	4 4		Table	2 con	it.		•		0000	25010
	9 903849	9.1.oct	2864995H	1 44		83	1	0 433776	4 00+	170000			
	9 903849	1.oct	5641431H	1 44 1 44	_	43	10	^U 433776	.4.oct	1726524 1724358	4H]	722	854
	9 903849	1.oct	1527073H	1 44	_	89	10	9 433776	.4.oct	2459364	104 101	722	796
	9 903849	.1.oct	5062263F6	46	_	45	10	433776	.4.oct	9384115	17 I	743	796
	9 903849	.1.oct	4172240H	1 44	•	91	10	433776	4.oct	5451747		744	796
	9 903849 9 903849	.1.oct	3959981H2	46		13 22	10	433776	4.oct	3603250		173 183	428
			4549428H1	45	27		10	433776.	4.oct	3571124	H1	185	485
			1527048H1	44	25		10		4.oct	6014182		192	494 422
			5062295H1	46	29		10 10		4.oct	1004309	R1	192	661
	9 903849. 9 903849.		256106H1	47	21		10		4.oct	19231251	H1 ·	194	459
	9 903849.		g1648330	47	20		10		4.oct	3969881	41 1	194	464
	9 903849.		g1638232	46	36	1	10	433776.4 433776.4	.oct	10043091	41 1	92	456
	903849		g519043 4550411H1	53	39	0	10	433776.4	.oct	g1678417	' 1	95	511
	9 903849.·		1777779H1	59	14	5	10	433776.4	1001	1504030H		99	456
	9 903849.1	Loct 3	3222580H1	57	34		10	433776.4	oct	g1940320	_	01	567
	10 433776.4	loct 5	033631H1	57	358		10	433776.4		g2556305 g2554138	_	10	433
	10 433776.4	loct n	1687856	469	737		10	433776.4		4763891H	2	12	626
	10 433776.4	oct 2	293262H1	472 471	854		10	433776.4	.oct	4266849H		26	483
	10 433776.4	oct 1	354926H1	471	711		10	433776.4	.oct	4350779H		28	348
	0 433776.4	.oct 4	465127H1	471	707 715		10	433776.4	oct	1724295T		31	332
	0 433776.4 0 433776.4	.oct 1;	354926F1	471	854		10	433776.4.	oct :	567093H1	26	77 30	798 405
			380506H1	474	773		10	433776.4.	oct (608247H1	. 27		425 529
1			90032H1	474	742		10 10	433776.4.	oct (607814H1	. 27		545
10			396456H1	474	736		10	433776.4.	oct (5093674H1	27		570
10			317480	496	864		10	433776.4.		939181H1	29	_	592
10			3675H1	496	796		10	433776.4.6 433776.4.6		006147H1	30		577
10	433776.4.		89927H1 3675T2	496	628		10	433776.4.0		427972H1		0 :	558
10	433776.4.		227083	496	821		10	433776.4.0	λCI] vct 1	923769H1			573
10) 433776.4.d	oct 64	266355	498	854		10	433776.4.0		923769R6 411986H1			727
10	433776.4.0	oct as	557641	507 508	796		10	433776.4.0		511411H1	322		552
10	433776.4.6	oct as	268469	510	796		10	433776.4.o	ct 2:	540387H1			579
10		oct g37	755408	510	796 796		10	433776.4.0	ct 6	078678H1	331 331		63
10 10		ct g2()47133	517	796		10	433776.4.0	ct 9:	34752H1	335		31 75
10			100966	517	796		10	433776.4.0	ct 93	34752R1	335	_	75 07
10			9730H1	519	794		10	433776.4.00	t 20	010868H1	334	_	32
10	433776.4.0 433776.4.0		300H1	537	767		10 10	433776.4.00	nt 93	34752T1	335	•	16
10	433776.4.00		87068	540	796		0	433776.4.00		73207H1	347	_	01
10	433776.4.00		37132	547	875		0	433776.4.00 433776.4.00		60027H1	349		11
10	433776.4.00		5017 32629	549	846			433776.4.oc		30254H1	355	60	05
10	433776,4,00	1 061		553 550	796	1	0	433776.4.oc		50377H1 847829	363	63	
10	433776,4,00	1 037		553 558	796	1	0	433776.4.oc		047829 23654H1	369	80)5
10	433776.4.00			573	796	1	0 4	433776.4.oc	122	23654T1	381	50)7
10	433776.4.oc	t a396			796 796	10	0 4	433776.4.oct	010	987932	380 380	81	
10 10	433776.4.oc	t g282	1439		796	10	0 4	133776.4.oct	527	75279H1	387	64 56	
10	433776.4.oc		96R1 !		796	10 10		33776.4.oct	186	7144H1	387	62	
10	433776.4.oct		96H1		796	10		33776.4.oct	g22	276753	399	75	
10	433776.4.oct			97	875	10		33776.4.oct	_	53602	402	85	
10	433776.4.oct			97	768	10		33776.4.oct 33776.4.oct		56942	403	85	
10	433776.4.oct				796	10	4	33776.4.oct	g35	96025	405	854	
10	433776.4.oct				796	10	•	33776.4.oct	807	740H1	406	592	2
10	433776.4.oct	1343			312	10	4	33776.4.oct	482	336H1 0231H1	406	575	
10	433776.4.oct	g1521		18 7	96	10	4	33776.4.oct	323	3233H1	406	595	
10	433776.4.oct	20089		29 7 29 8	96	10	43	33776.4.oct	1616	5991T6	413	666	
10	433776.4.oct	16169	56T6 63		11	10	43	33776.4.oct	g373	34566	416	816	
10	433776.4.oct	g4333	263 62		96 96	10	43	33776.4.oct	g408	88877	416 419	857	
10	433776.4.oct	41735	46H1 64		96 97	10	43	33776.4.oct	g406	9104	420	857	
10 10	433776.4.oct	g4270	379 64	_	57 63	10	43	3776.4.oct	1493	172H1	431	854 660	
10	433776.4.oct	g4450	858 ₆₅		9 6	10	43	3776.4. ct	2005	329H1	431	669 619	
10	433776.4.oct 433776.4. ct	222969	90H1 65		58	10 10	43	3776.4.oct	g184	7522	437	852	
10	433776.4. ct	63874		1 94		10	43	3776.4.oct	g342	5153	441	869	
	433776.4.oct	g34309		7 79		10	43	3776.4.oct	g121	0346 .	443	856	
	433776.4.oct	235266 g30528		0 79	6	10	43	3776.4.oct	g3679	9055	446	849	
		900028	21 719	9 79	6	10	433	3776.4.oct	40194		448	733	
						67			7013	137H1 4	149	704	
					,	- ,	•						

				Table	2 cont.		1 0 1/ 0 300/ 23010		
13	336430.2.dec	g2568596	946	1420	14	242269.2.dec	g3446472	8	341
13	336430.2.d c	5327737H1	952	1199	14	242269.2.dec	g1300544	6	304
13	336430.2.dec		952	1185	14	242269.2.dec	g1300552	6	191
13	336430.2.dec	•	958	1421	14	242269.2.dec	•	8	186
13	336430.2.dec	•	963	1421	14	242269.2.dec	0	8	459
13	336430.2.dec	•	963	. 1421	14	242269.2.dec		9	472
13 13	336430.2.dec 336430.2.dec		966	1422	14	242269.2.dec	•	9	410
13	336430.2.dec		967 980	1425 1433	14 14	242269.2.dec 242269.2.dec	93755968	8	392
13	336430.2.dec	g2094705	982	1433	14	242269.2.dec	J	8	359
13	336430.2.dec	O		1424	14	242269.2.dec	1351344F1	8 12	208 · 378
13	336430.2.dec	0	1007	1421	14	242269.2.dec		12	61
13	336430.2.dec	1696181H1	1015	1229	14	242269.2.dec	410843R1	12	501
13	336430.2.dec	g4088599	1030	1421	14	242269.2.dec		12	218
13	336430.2.dec	•	1032	1430	14	242269.2.dec	414883H1	12	185
13	336430.2.dec		1032	1436	14	242269.2.dec		12	184
13 13	336430.2.dec	•	1036	1421	14	242269.2.dec		12	171
13	336430.2.dec 336430.2.dec		1043 1045	1430	14	242269.2.dec		12	171
13	336430.2.dec		1045	1422 1427	14 14	242269.2.dec 242269.2.dec	412381H1	12	164
13	336430.2.dec		1065	1403	14	242269.2.dec	855180H1 g5101470	12 12	145 410
13	336430.2.dec		1085	1423	14	242269.2.dec		12	297
13	336430.2.dec		1086	1425	14	242269.2.dec		67	430
13	336430.2.dec	g3086789	1087	1426	14	242269.2.dec	g5232210	72	521
13	336430.2.dec	-	1100	1422	14	242269.2.dec	g2569754	74	507
13	336430.2.dec	~	1105	1431	14	242269.2.dec	g4074654	75	485
13	336430.2.dec	•	1107	1422	14	242269.2.dec		75	389
13 13	336430.2.dec 336430.2.dec	g752027 g2905252	1127 1136	1424	14	242269.2.dec		75	348
13	336430.2.dec		1159	1421 1423	14 14	242269.2.dec 242269.2.dec	g1202202 118555T6	75 84	239
13	336430.2.dec	•	1165	1421	14	242269.2.dec	6614443H1	84 84	585 583
13	336430.2.dec		1166	1421	14	242269.2.dec	1222034H1	88	258
13	336430.2.dec	g3172766	1206	1422	14	242269.2.dec	1222034T1	88	258
13	336430.2.dec	g2809576	1212	1421	14	242269.2.dec	4032139H1	88	213
13	336430.2.dec		1227	1427	14	242269.2.dec	1560349T6	88	191
13	336430.2.dec	g2807042	1233	1425	14	242269.2.dec	942024T1	88	184
13	336430.2.dec	g3431342	1242	1421	14	242269.2.dec	J	88	350
13 13	336430.2.dec 336430.2.dec	g3431049 g3233026	1267	1421	. 14	242269.2.dec		105	338
13	336430.2.dec		1272 1273	1421 1411	14 14	242269.2.dec 242269.2.dec	1351344H1	126	378
13	336430.2.dec		1285	1421	14	242269.2.dec	1565904H1 5812349H1	128 146	349 472
13	336430.2.dec	g3092413	1291	1425	14	242269.2.dec	5328191H1	156	395
13	336430.2.dec	1559410T6	1301	1379	14	242269.2.dec	3515332H1	160	417
14	242269.2.dec	g1242484	1	297	14	242269.2.dec	3787803H1	173	279
14	242269.2.dec		1	458	14	242269.2.dec	5531343H1	229	471
14	242269.2.dec	g3017107	1	274	14	242269.2.dec	5979339H1	234	533
14	242269.2.dec	g2716604	1	77	14	242269.2.dec		240	303
14 14	242269.2.dec 242269.2.dec	g3539600	1	355	14	242269.2.dec		310	524
14	242269.2.dec	g5528437 1560349H1	1 2	281 184	14 14	242269.2.dec	1756754R6	344	612
14	242269.2.dec	q1281302	2	415	14	242269.2.dec 242269.2.dec		365 366	952 672
14	242269.2.dec	g3182199	3	245	14	242269.2.dec		366 390	672 660
14	242269.2.dec	g4070944	3	408	14	242269.2.dec	1671523H1	392	587
14	242269.2.dec	g5235258	5 ·	465	14	242269.2.dec		396	676
14	242269.2.dec	g5232045	5	346	14	242269.2.dec		394	661
14	242269.2.dec	g3679020	5	382	14	242269.2.dec	5913540H1	420	699
14	242269.2.dec	1351344F6	8	378	14	242269.2.dec	1456573H1	430	708
14	242269.2.dec	g5101469	7	411	14	242269.2.dec		443	689
14 14	242269.2.dec 242269.2.dec	g2167296	5	353 457	14	242269.2.dec		450	675
14	242269.2.dec	g5232575 942024H1	8 8	457 184	14	242269.2.dec		454	1030
14	242269.2.dec	942024R1	8	184	14 14	242269.2.dec 242269.2.dec	g1317243 g1317236	476 505	894 805
14	242269.2.dec	3127564H1	8	141	14	242269.2.dec	•	505 512	895 773
14	242269.2.dec	g4436066	. 8	454	14	242269.2.dec		582	830
14	242269.2.dec	g4850437	8	398	14	242269.2.dec	1525743H1	665	810
14	242269.2.dec	g2740407	8	391	14	242269.2.dec		764	874
14	242269.2.dec	g3231712	8	362	14	242269.2.dec	4669147H1	768	1016
14	242269.2.dec	g3427933	8	<u>3</u> 50	14	242269.2.dec	411214H1	1	241
					70	•			

40		,		T	able 2	CORt			1	PCT/	US00/25610
12	234828.6.oct		39	300							, == 0.20
12	234828.6.oct	990620H1	37	332		12	234828	.6.oct	267060	5U4	
12.	234828.6.oct	2857770H1	39			12	234828	.6.oct	338536		25 307
12	234828.6.oct	2456375H1	39	295		12	234828	6.oct	132474		27 295
12	234828.6.oct	1452171H1	40	279		12	234828.	6.oct			27 277
12	234828.6.oct	2443307H1		294		12	234828.	6.00		9H1	26 130
12	234828.6.oct	2477496H1	- 40	280		12	234828.	6.000	6176386	5H1	27 316
12	234828.6.oct	729737H1	40	270		12	234828.	0.0Ct	1502616	5H1	14 256
12 2	234828.6.oct	729737R1	41	270		12	234828.	0.001	3591749	9H1 2	27 329
12 2	234828.6.oct	729/3/H1	41	381		12	224920	9.0CT	2443723	BH1 2	7 253
12	234828.6.oct	3482624H1	41	193		12	234828.6	o.oct	3204501	H1 2	8 153
12 2	34828.6.oct	3199337H1	45.	309		12	234828.6	o.oct	2015825	H1 2	8 327
12 2	34838 C	1340368H1	45	222		12	234828.6	oct.	2733437	H1 2	
	34828.6.oct	4265872H1	45	242			234828.6	.oct	2198080	H1 2	
	34828.6.oct	3871172H1	46	318		12	234828.6	.oct	14449481	H1 2	,
12 2	34828.6.oct	2202960H1	46	284		12	234828.6	.oct	2961687	H1 2	
	34828.6.oct	1839888H1	46			12	234828.6	oct	61799911		
12 2	34020.6.0Ct	1823204H1	46	286		12	234828.6	oct	2198495H		
12 23	34028.6.0ct	2740138H1	53	263		12	234828.6.	oct	2906122H		
12 23	74028.6.0ct	5488475H1	71	295		12	234828.6.	oct	3297819H	11 29	
12 23	74028.6.0ct	5119401H1		321		12	234828.6.		20012121	_	
12 23	34828.6.oct /	6158348H1	101	377	1	12 :	234828.6.		2901213H	11 30	
12 23	4828.6.oct 6	g1967590	139	414	1	12 2	234828.6.		2782627H	1 29	279
12 23	4828.6.oct	2345622H1	153	615	1	12 2	234828.6.		1625443H		268
12 23	4828.6.oct 2	2241354H1	155	403	1	2 2	234828.6.0		6178209H	1 30	303
12 23.	4828.6.oct 3	3687713H1	171	421			336430.2.0		3208193H	1 29	297
12 23		895482H1	171	478			36430.2.0		3593730H	1 1	168
12 234		469071H1	172	431		_	36430.2.d		2781490H	1 1	242
12 234	1828.6.oct 5	469000111	174	421	1:		36430.2.0		3882669H	1 7	296
12 234	1828.6.oct 1	468888H1	174	440	1:		36430.2.d		3520838H1	1 19	347
12 234		453408H1	177	391			36430.2.d		3574135H1	1 10	146
			213	459	13		36430.2.d		559410H1	20	247
		08399H1	225	443	13	3 3	36430.2.d	ec 1	559410F6	20	229
		473672H1		462	13		36430.2.d	ec 2	266238H1	29	181
		358095H1	~ 4 ~	465	13) <u>(</u>	36430.2.de	ec 1:	299583H1	31	227
		300564H1		513			36430.2.de	3C 31	045047H1	34	
	828.6.oct 46	550680H1		528	13	33	86430.2.de	ec 11	227115H1	35	164
	920.6.0Ct 17	'53653H1 ·	`	471	13		6430.2.de	C 15	535201H1	45	277
12 2348	528.6.oct 17	51064H1		155	13	33	6430.2 de	C 01	1300630		257
	28.6.oct 16	13647H1		133 121	13	33	6430.2.de	C 02	2111158	105	520
12 2348	28.6.0ct 14	00857H1 2	'		13	33	6430.2.de	C 23	87085H1	157	360
12 2348	28.6.oct 13	38695H1 2		36	13	33(6430.2.de	C 02	210797	240	425
12 2348	<0.6.oct 887	74 701		75 20	13	336	6430.2.de	c 37	61768H1	269	725
12 2348	28.6.oct 21r	3040011	`	36	13	336	6430.2.de		44503H1	281	583
12 2348	28.6.oct 266	20000114		6 0	13	336	3430.2.ded	: 40	62190H1	305	802
12 23482	28.6.oct 755			63	13	336	3430.2.dec		62028H1	344	512
12 23482	28.6.oct 711	700	~ .	91	13	336	430.2.dec		409916	396	898
12 23482	28.6.0ct 362	2670114		38	13	336	430.2.dec		41464H1	413	726
12 23482	28.6.oct 616		53 60	_	13	336	430.2.dec		+1404M1	423	681
12 23482	8.6.oct 430	004011			13	336	430.2.dec		71874H1	452	533
12 23482	8.6.oct 430		•		13	336	430.2.dec		37615H1	486	726
12 23482	8.6.oct 362	070044			13	336	430.2.dec	400	0922H1	511	678
12 23482	8.6.oct 5881	40= 44			13	3364	430.2.dec		1269F6	603	1033
12 23482	8.6.oct 5889	\4==\.			13	3364	130.2.dec		1269H1	603	860
12 234821					13	3364	130.2.dec	510	6212H1	659	921
12 234828				8	13	3364	130.2.dec	g209	94638	671	1091
12 234828		432F6 40		1	13	2264	130.2.dec	g209	94369	676	1077
12 234828	36 oot 0450	763H1 406	615		13	3364	30.2.dec	1688	8367F6	681	1229
12 234828		375T6 406	916		13	3364	30.2.dec	1688	3376H1	681	916
12 234828		432H1 408	520			3364	30.2.dec	1688	3985H1	681	
12 234828		963H1 407	684		13	3364	30.2.dec	4003	3277H1	714	895 978
12 234828		868H1 1	261		13	3364	30.2.dec	g189	1651	735	9/8
	.6.oct 35386	688H1 12	279		13	33643	30.2.dec	5604	707H1	738	1123
	.6.oct 24406	543H1 14	266		13	33643	30.2.dec	5369			1013
	.6.oct 15026	80H1 1A			13	33643	30.2.dec	5370	00111	741	976
12 234828.	.b.oct 29728	304H2 15	302		13	33643	30.2.dec	6123	400		898
12 234828.	6.oct 15082	18H1 20	316		13	33643	0.2.dec	18001			1347
12 234828.	6.oct 322na	62H1 25	194		13	33643	0.2.dec	6300			1009
12 234828.6	6.0ct 14000	A 41 4	314		13	33643		シンフェイ	30044	786	1379
12 234828.0	6.oct 31486	04114	245			33643		38いこっ		797	1015
12 234828.6	6.oct 61865	C1 14	294		13	33643		1011		830 -	108
	3.000	bH1 25	295		13	33643	0.2.dec 3	10112		330 ·	379
			-	69	`			39988	פייאאון פ		230
				09	,						

18	235983.6.dec	g390184	2995	3311	18	235983.6.dec	2465140H1	3782	4006
18	235983.6.d c	4220392H1	3005	3279	18	235983.6.dec	q3174357	3788	4082
18	235983.6.dec	3286977H1	3012	3260	18	235983.6.d c	4024334H1	3790	4081
		g616559	3020	3327	18				
18	235983.6.dec	-				235983.6.dec	1499362H1	3787	3978
18	235983.6.dec	2192319H1	3033	3264	18	235983.6.dec	4646510H1	3789	4056
18	235983.6.dec	5683292H1	3051	3287	18	235983.6.d c	1594535H1	3789	3999
18	235983.6.dec	4182518H1	3054	3332	18	235983.6.dec	898100H1	3797	4054
18	235983.6.dec	926283H1	3062	3334	18	235983.6.dec	4837242H1	3798	4012
18	235983.6.dec	3491246H1	3067	3343	18	235983.6.dec	898100R1	3797	4374
18	235983.6.dec	4601526H1	3068	3332	18	235983.6.dec	4837492H1	3799	4086
18	235983.6.dec	q883924	3070	3132	18	235983.6.dec	3573408H1	3799	4096
18	235983.6.dec	4898255H1	3078	3326	18	235983.6.dec	4837274H1	3799	4053
							_		
18	235983.6.dec	g612635	3095	3379	18	235983.6.dec	g1391806	3807	4225
18	235983.6.dec	1501609H1	3091	3287	18	235983.6.dec	g4901871	3806	4225
18	235983.6.dec	5435484H1	3095	3275	18	235983.6.dec	5861440H1	3806	4093
18	235983.6.dec	1710332H1	3109	3332	18	235983.6.dec	5152105H1	3809	4084
18	235983.6.dec	3706933H1	3120	3405	18	235983.6.dec	833965H1	3809	4080
18	235983.6.dec	991301H1	3125	3430	18	235983.6.dec	g2331330	3810	4124
18	235983.6.dec	991301R1	3125	3561	18	235983.6.dec	2053601H1	3815	4070
18	235983.6.dec	4670213H1	3130	3378	18	235983.6.dec	903670H1	3815	4007
18	235983.6.dec	1815401T6	3640	4185	18	235983.6.dec	2875944H1	3816	4091
18	235983.6.dec	5610495H1	3639	3898	18	235983.6.dec	4402735H1	3816	4074
18	235983.6.dec	g1050006	3647	3981	18	235983.6.dec	5530194H1	3820	4068
18	235983.6.dec	g711179	3646	3848	18	235983.6.dec	g2752396	3818	4234
18	235983.6.dec	1317687H1	3646	3820	18	235983.6.dec	6009383H1	3826	4108
18	235983.6.dec	1968426H1	3656	3917	18	235983.6.dec	g2873872	3825	4228 -
18	235983.6.dec		3658	4226	18	235983.6.dec	3658872H1	3833	4112
18	235983.6.dec	4933278H1	3657	3929	18	235983.6.dec	g1153144	3831	4237
18	235983.6.dec	4110987H1	3658	3755	18	235983.6.dec	2500844H1	3842	4072
18	235983.6.dec	878460H1	3658	3909	18	235983.6.dec	6312876H1	3844	4372
18	235983.6.dec	2195022H1	3661	3903	18	235983.6.dec	1428214T6	3849	4185
18	235983.6.dec	4152080H1	3664	3925	18	235983.6.dec	2244834H1	3850	4105
18	235983.6.dec	984860R1	3667	4117	18	235983.6.dec	g3869186	1	4539
18	235983.6.dec	984860H1	3667	3888	18	235983.6.dec	3699775H1	61	346
18	`235983.6.dec	3573936H1	3673	3955	18	235983.6.dec	5047477H1	147	377
18	235983.6.dec	3088228H1	3677	3952	18	235983.6.dec	3381586H1	. 145	394
					18			146	679
18	235983.6.dec	1781737H1	3680	3878		235983.6.dec	6484838H1		
18	235983.6.dec	g2053250	3686	4062	18	235983.6.dec	3584956H1	147	464
18	235983.6.dec	4254583H1	3687	3941	18	235983.6.dec	6476413H1	148	678
18	235983.6.dec	g1101456	3688	3907	18	235983.6.dec	5047477F6	147	671
18	235983.6.dec	g982339	3689	4031	18	235983.6.dec	6476530H1	148	663
18		g1042775		3974	18			147	346
	235983.6.dec	•	3689			235983.6.dec	5047423H1		
18	235983.6.dec	6350466H2	3692	4013	18	235983.6.dec	3586619H1	147	341
18	235983.6.dec	3624071H1	3692	3876	18	235983.6.dec	3073571H1	147	432
18	235983.6.dec	4425320H1	3694	3970	18	235983.6.dec	353081H1	154	378
18	235983.6.dec	4793731H1	3698	3978	18	235983.6.dec	3288503H1	155	418
			3699	3981	18	235983.6.dec			
18	235983.6.dec	4991525H1					121178H1	160	232
18	235983.6.dec	2346595H1	3699	3952	18	235983.6.dec	g2703916	489	726
18	235983.6.dec	2666581H1	3699	3942	18	235983.6.dec	3119591H1	505	773
18	235983.6.dec	3203383H1	3702	3961	18	235983.6.dec	g4104518	611	2596
18	235983.6.dec	g1046477	3719	4063	18	235983.6.dec	g1471244	675	725
18	235983.6.dec	g714049	3719	4027	18	235983.6.dec	4994487H1	. 916	1180
18	235983.6.dec	5471836H1	3725	3984	18	235983.6.dec	3353480H2	953	1127
18	235983.6.dec	3692694H1	3739	4020	18	235983.6.dec	1477782H1	1030	1238
18	235983.6.dec	3345345H1	3741	3983	18	235983.6.dec	5950615H1	1047	1298
18	235983.6.dec	2530908H1	3749	3999	18	235983.6.dec	2525583F7	1097	1544
18	235983.6.dec	1446674H1	3760	4024	18	235983.6.dec	2525583H1	1097	1343
18	235983.6.dec	1397742H1	3761	4023	18	235983.6.dec	434615H1	1122	1347
18	235983.6.dec	1400409H1	3761	3975	18	235983.6.dec	5153342H1	1281	1547
18	235983.6.dec	g1141309	3767	4119	18	235983.6.dec	6281520H1	1337	1611
		•							
18	235983.6.dec	3916312H1	3768	4052	18	235983.6.dec	531966H1	1353	1596
18	235983.6.dec	4598159H1	3771	4029	18	235983.6.dec	2608473F6	1364	1818
18	235983.6.dec	855084H1	3777	4024	18	235983.6.d c	2608473H1	1364	1582
18	235983.6.dec	855084R1	3777	4168	18	235983.6.dec	6282763H1	1367	1611
18	235983.6.dec	382681H1	3777	3922	18	235983.6.dec	6289464H1	1407	1611
18	235983.6.dec	4511379H1	3779	4009	18	235983.6.d c	4997470H1	1419	1674
18	235983.6.dec	g2806189	3780	4225	18	235983.6.dec	6006170H1	1421	1697
18	235983.6.dec	1964217H1	3782	4051	18	235983.6.dec	6285201H1	1430	1611

110 01/25558			
,,	77		PCT/IICOO to a
15 432120.2.d c 6245865H	. 1	able 2 cont.	PCT/US00/25610
		17 460295 5 dos	
	1 1 511	17 400005.5.dec 933671	88 56 521
	161 593	17 400295.5.dec g408010	00 56 454
	212 500	17 400295.5.dec g472919	90 56 979
	1 1 200	'' 400495.5.dec a40000	No. 50 3/3
16 100000.0.dec 1669868F6	1 405	**************************************	17 = 20/
16 10000.U.UEC 1668592H1	1 227	" TOUZSO.5.dec asaga ₄₆	4
16 100000.0.dec 1669868H1	1 225	460295.5.dec 319502L	34
16 190000.6.dec g2163372	12 479	460295.5 dec gaggego	
198000.6.dec 4217360H1		460295.5.dec 0317320	6 444
190000.6.dec 3115007H1		17 460295.5.dec 4200050	LI4 430
130000.6 dec 2206074114		10 235983.6 dec 40200cm	
19000U.B.Dec 582217U4			
198060.6.dec 631476014	,		11 2138 220₄
198060.6.dec 207456714	22 542		11 2157 2447
198060.6.dec 250224014	22 269		11 2170 0440
198060.6.dec 164691014	22 260	10 20000.0.UEC 35/6941H	11 2186 2450
198060,6,dec 75400704	24 225		6 2184 2050
198060.6.dec 754007114	31 540		1 2184 2400
198060.6.dec 1216707114	31 230	10 352928H1	2204 2400
198060.6 dec 163670044	26 268	10 ====================================	1 2225 2470
	26 242	235983.6.dec 5065937H	1 2227 2420
	28 340	19 233983.6.dec 5545038H	1 222
	29 463	235963.6.dec 370706H1	2249 2464
	29 332	235983.6.dec 4342101H1	2253 2504
	30 272	200503.b.dec 3115000112	
	31 270	200300.D.DEC 1676040F0	070
16 10000.0.dec 3521542H1	35 304	200903.b.dec 167504014	
16 10000.0.dec 92154340	34 494	200300,0.dec 01707040	
16 10000.0.dec 92055182	34 424	235983.6.dec 2224910114	2000
16 3699676H1	34 290	200300.D.DRC 4171460114	2355 2597
16 1000-0-0-0-0-0-0-16/35H1	35 303	235983.6.dec 01939937	2414 2677
16 130000.6.dec 2078062H1	34 324	235983.6.dec 183400U4	2426 2907
16 100000.0.dec 92240557	39 394	18 235983.6.dec 01812050	2478 2651
16 198060.6.dec 6168723H1	52 377	16 235983.6.dec 3050415114	2489 2872
16 155225H1	75 338	18 235983.6.dec d2002020	2505 2644
196000.6.dec 1395663H1		16 235983,6,dec 04152315	2512 2877
190000.b.dec 5047207114		18 235983.6.dec 211521414	2525 2928
16 190000.6.dec 1615709F6		16 235983.6.dec 01301010	2522 2802
190000,6.dec 1615700U4			2545 2914
190000.b.dec 1615650114		18 235983.6.dec g1492984 235983.6.dec 5616747H1	2545 275 7
198060,6,dec 612141514			2676 2965
198060,6,dec 497605514	552		2686 2978
198060.6.dec 5571120U4	09 384		2701 2963
198060.6.dec 50162014	36 338		2747 3032
198060.6.dec 02505782	48 407		2763 3040
10 198060.6.dec 384305044	53 550	1p 92002465	2773 3189
198060.6.dec 287604114	55 466		2778 3084
198060,6,dec 464217014	63 435		2776 3038
198060.6.dec d2251550	67 432	19 20500.0.060 4935114H1	2782 2880
198060.6.dec 4307774114	35 494	10 005000.0.dec g//2632	2784 3120
198060,6 dec 1971040F0			2804 3005
198060.6.dec 197104014		10 20000.0.dec 1502463H1	2811 3087
198060 6 dec 41000=111 23		19 005000.0.dec 1502565H1	2811 3078
198060 6 dec 10007007	4 528	19 235983.6.dec g4152317	2817 3146
198060 6 dos 1000-	8 613	19 235983.6.dec 2319441H1	
16 10000 1000/93H1 24	8 502	19 235983.6.dec 188577H1	
	5 479	233983.6.dec 476050U4	2000
16 100000.0.dec 1527256H1 260		235983.6.dec g2053717	
16 1000.0.dec 152/264H1 260	474	200983.6.dec 1640200114	
16 100000.0.dec 5884461H1 286		18 235983.6.dec 476070014	2901 3100
16 100000.0.dec g5639170 373		16 235983.6.dec 472140214	2907 3195
16 100000.0.dec 4913811H1 437		18 235983.6.dec 472122414	2907 3186
16 198000.6.dec 4409109H1 410		18 235983.6.dec 2667214U4	2907 3165
16 196000.6.dec 4611874H1 480		18 235983.6.dec 181540150	2954 3199
16 196060.6.d C 232531H1 514		18 235983.6.d c 181540414	956 3354
16 196060.6.d c 2829981H1 516	670 700	18 235983.6.dec 120215014	956 3214
16 198000.b.d c 4795943H1 663	786	10 235983.6.dec 120210414	956 3209
16 198060.6.d c 3340413H1 660		16 235983.6.dec 215527014	956 3210
1/ 460295.5.d c 621446014	917	10 235983 6 dec 207075444	958 3206
1	520	10 235983.6 dec 305540744	966 3258
•	. 71		979 3261
	/1		

Table 2 cont. 18 235983.6.dec 4301987H1 4164 4405 18 235983.6.dec g718586 4278 4517 18 235983.6.dec g1773776 4165 4409 18 235983.6.dec g2445128 4285 4534 18 235983.6.dec 3419076T6 4197 4508 18 235983.6.dec 92053618 4285 4537 18 235983.6.dec g1264115 4202 4539 18 235983.6.dec 2563295H1 4289 4541 18 235983.6.dec 928449T6 4210 4501 235983.6.dec 18 g1049755 4299 4546 235983.6.d c 18 q4874751 4210 4545 18 235983.6.dec g779371 4300 4545 18 235983.6.dec 2005565H1 4213 4413 18 235983.6.dec 1979178H1 4299 4539 18 g1148504 235983.6.dec 4215 4539 18 235983.6.dec g3871331 4305 4539 18 235983.6.dec g866362 4219 4545 18 235983.6.dec 910538H1 4312 4533 18 235983.6.dec 3956647H1 4219 4508 18 235983.6.dec 549362F1 4335 4541 18 235983.6.dec 4359163H1 4220 4447 18 235983.6.dec 2275266H1 4338 4539 18 235983.6.dec g883309 4224 4567 18 235983.6.dec 549362H1 4346 4541 18 235983.6.dec 4223 g723173 4525 18 235983.6.dec 3808941H1 4346 4533 18 235983.6.dec g3840912 4225 4544 18 235983.6.dec q2589294 4349 4547 18 235983.6.dec q2986196 4228 4541 18 235983.6.dec 287095H1 4351 4533 18 g2715739 235983.6.dec 4227 4536 18 235983.6.dec 2662364F6 4364 4533 18 g1046478 235983.6.dec 4227 4507 18 235983.6.dec 2662364H1 4364 4533 18 235983.6.dec g4088760 4230 4533 18 235983.6.dec 5847389H1 4366 4533 18 235983.6.dec g3425690 4230 4533 18 235983.6.dec 5872270H1 4369 4441 18 235983.6.dec 4466803H1 4231 4383 18 235983.6.dec 4298534H1 4372 4539 18 235983.6.dec 767529H1 4236 4471 18 235983.6.dec g1210948 4383 4542 18 235983.6.dec g612988 4237 4541 18 235983.6.dec a1137312 4397 4541 18 235983.6.dec g2817035 4237 4539 18 235983.6.dec 811100T1 4419 4497 18 235983.6.dec 2430718H1 4239 4477 18 235983.6.dec 811100H1 4419 4527 18 235983.6.dec g5510981 4243 4554 18 235983.6.dec g2270136 4429 4534 18 235983.6.dec g3279191 4243 4545 18 235983.6.dec 233542H1 4430 4533 18 235983.6.dec 3040021H1 4243 4450 18 235983.6.dec g982292 4450 4512 18 235983.6.dec 219012H1 4243 4405 18 235983.6.dec g1792123 4452 4539 18 235983.6.dec 5566988H1 4243 4376 18 235983.6.dec g3057972 4466 4542 18 235983.6.dec g1860203 4243 4544 18 235983.6.dec g3839889 4479 4548 18 235983.6.dec g3769996 4243 4543 18 235983.6.dec 4467167H1 4479 4533 18 235983.6.dec g3057160 4243 4543 18 235983.6.dec g1691482 4487 4543 18 235983.6.dec g2206143 4243 4547 18 235983.6.dec 2942208H2 3134 3398 g3231269 18 235983.6.dec 4243 4546 18 235983.6.dec 3500378H1 3137 3430 18 235983.6.dec g3933929 4243 4541 18 235983.6.dec 3873691H1 3143 3409 18 235983.6.dec g4152314 4243 4541 18 235983.6.dec 4644974H1 3144 3402 18 235983.6.dec g3917367 4243 4539 18 235983.6.dec 3620275H1 3158 3412 18 235983.6.dec g3331035 4243 4539 18 235983.6.dec 5198954H1 3158 3327 18 235983.6.dec g2741042 4243 4540 18 235983.6.dec 6412923H1 3158 3514 18 g5541034 235983.6.dec 4243 4538 18 235983.6.dec 4692566H1 3168 3408 18 235983.6.dec g3675471 4243 4538 18 235983.6.dec 4941568H1 3169 3437 18 235983.6.dec g3091778 4243 4533 18 235983.6.dec 3680473H1 3185 3467 18 235983.6.dec q3203015 4243 4532 18 235983.6.dec 3687818H1 3194 3489 18 235983.6.dec 1565654H1 4243 4386 18 235983.6.dec 5865911H1 3196 3467 18 235983.6.dec g1379563 4243 4533 18 235983.6.dec 5676145H1 3221 3457 18 235983.6.dec g3191370 4243 4532 18 235983.6.dec g1691481 3230 3594 18 235983.6.dec g3233027 4243 4533 18 235983.6.dec 2225035H1 3254 3490 18 235983.6.dec g29057 4251 4539 18 235983.6.dec 157436H1 3265 3476 18 235983.6.dec q5108814 4249 4535 18 235983.6.dec 157436R1 3267 3781 18 235983.6.dec 2126663H1 4254 4523 18 235983.6.dec q2069871 3277 3638 18 4254 235983.6.dec g1101555 4533 18 235983.6.dec 4445787H1 3515 3278 18 235983.6.dec 1677718H1 4257 4498 18 235983.6.dec 3616513H1 3310 3614 18 235983.6.dec q877366 4257 4541 235983.6.dec 18 158334H1 3315 3482 235983.6.dec 18 q842556 4256 4533 18 235983.6.dec 726102H1 3614 3838 18 235983.6.dec q1011570 4258 4546 18 235983.6.dec 1319351H1 3319 3542 18 235983.6.dec g1039989 4267 4542 18 235983.6.dec 4447125H1 3349 3621 18 235983.6.dec g2821823 4265 4549 18 235983.6.dec 4648142H1 3358 3618 18 235983.6.dec q3116989 4265 4539 18 235983.6.dec 3895241H1 3361 3524 18 235983.6.dec g782426 4269 4547 18 235983.6.dec 2022857H1 3364 3589 18 235983.6.dec g1939717 4269 4546 18 235983.6.dec 5013742H1 3362 3632 4522 18 235983.6.dec q2841416 4267 18 235983.6.dec 3407443H1 3369 3605 18 235983.6.d c g4511390 4270 4533 18 235983.6.dec 1427630F6 3380 3949 18 235983.6.dec g2821824 4280 4553 18 235983.6.dec 1427630H1 3380 3603 18 235983.6.dec 536291H1 4274 4521 18 235983.6.dec 4890434H1 3388 3568 18 235983.6.dec g3802343 4276 4541 18 235983.6.dec g705111 3390 3451 18 744021R1 235983.6.dec 4277 4533 18 235983.6.dec 5301717H1 3420 3637 18 235983.6.dec 744021H1 4277 4517 18 235983.6.dec 1831110H1 3420 3670 18 235983.6.dec g1727350 4280. 4544 18 235983.6.dec 1220017H1 3429 3658

10	Tak	ole 2 cont.	PCT/US00/25610
18 235983.6.dec 3617263H1		ne 2 cont.	27.0300/23010
235983.6.d C 6292515H1	4444	18 235983.6.dec	1010740
233983.6.d c 6289496H1	444	235983.6.dec	1919746H1 3982 4225
255963.6.dec 5636916H1	.011	16 235983,6,dec	63490514 3982 4217
19 200903.0.0 C 6355532H1		10 235983.6.d c	3982 4217
18 233963.6.dec 4070803H1	4.45	10 235983.6.dec	3986 4423
18 235983.6.dec 4153164H1		¹⁶ 235983.6.dec	3991 4510
18 203963.6.dec g2810741	1500	16 235983.6.dec	20009/8H1 3990 4217
235983.6.dec g2901231	1500	¹⁶ 235983.6.dec	2554512H1 3990 4217
19 235983.6.dec g3050079	150-	⁷⁶ 235983.6 dec	167504075 3996 4217
233983.6.dec 6286265H2	101-		1675942T6 4013 4495
200300.D.DRC 628/160114	4040		5077440HP 4015 4225
233963.6.dec 6289955H1			5077446H2 4015 4304
19 255963.6.dec 6290838H1			24242411 4014 4217
18 233963.6.dec 6290954H1	1010	¹⁰ 235983.6.dec 1	701405111 4020 4203
10 233963.6.dec 3613944H1		10 235983.6 dec 3	701425H1 4030 4223
200963.6.dec 3154382H1	472		1400079 4050 4340
200303.6.dec 3363928H1	1700	10 235983.6.dec 4	74244514 4053 4225
2005903.0.dec 2205748H1	4777	235983.6.dec p	42415H1 4054 4306
18 200503.6.dec 2205748F6		10 235983.6 dec 91	16546T1 4056 4546 16546H1 4056 4344
18 203903.0.dec g395643		10 235983.6.dec as	ACCEA!
235983.6.dec 3075529F6	00	√ 16 235983.6.dec 44	4060 4542
19 233963.b.dec 3419076F6	1010	10 235983.6.dec 10	106000111 4066 4321
19 205505.0.dec 5668869H1		16 235983.6.dec 30	1400014
18 205903.0.dec 4670760H1 1		18 235983.6.dec at	27050
19 203903.0.dec 928449H1 1	849 2104 870 2132	10 235983.6 dec 31	5004075
235983.6.dec 928449R1 1		235983.6.dec 46	
19 200903.b.dec 928449R6 1		235983.6.dec a20	154055
19 233963.6.dec 3492637H1 1	871 2323 918 2181	10 235983.6.dec 054	120445
19 200903.0.dec 484275R6 10	990 2490	10 235983,6,dec dae	01079
18 205903.6.dec 484275H1 16	990 2231	235983.6.dec 054	25054
18 235000 g1312232 3	351 4278	235983.6.dec 301	EC70.11
18 233963.6.dec g870634 36	356 4234	10 235983.6.dec 357	F200114
18 005005.0ec 841880R1 38	366 4317	10 235983.6.dec 264	580001
10 0050-10.0ec 598558H1 38	62 4092	10 235983.6 dec -40	00642
18 205050.0.dec 607754H1 38	65 4120	10 235983.6.dec 636	14504
18 305000 04 1880H1 38		10 235983.6.dec dage	MAEN: 1000
18 335000.0.dec g2139402 38		10 235983.6.dec 0456	18272
10 005-0-0.0.000 92205778 30		10 235983.6.dec 0567	4002
19 00500.0.dec 92148308 386		10 235983.6.dec d343	3200 4000
19 20500.0.000 9/81601 390		10 235983.6.dec 0256	844E 4000
	6 4008	10 235983.6.dec 3551	2011
18 205000.000 94152316 380	9 4215	200900.0.dec 3506/	CIOTA TEST
19 005000 15/436F1 300			7570
16 225000 0	6 4121	TOUROU.D. GEC 04224	1047
16 235983 6 dog 44975	7 4093	200963,6,dec 01070	1154 1104 4540
	1500	235983.6.dec g4649	188 4110 4500
	1 1001	235983.6.dec g3959	037 4124 4505
16 235983 6 dos 4048-111 3925	4139	18 235393.6.dec g3693	698 4124 4530
10 235983.6 dec 252550070 3924	4496	200903.6.dec g4568	110 4124 4500
18 235983.6.dec 100024070 3931	4495 1	g 225000.0.dec 943923	393 4125 4544
16 235983.6.dec 3075500To	4495 1	255983.b.dec g40883	342 4127 4545
16 235983.6.dec 260047075 3540	4509 ₁	235963.6.dec 422198	37H1 4126 4404
16 235983.6 dec 6115700111 3947	4500	200903.b.dec 422139	3H1 4126 4448
18 235983.6 dec 2007449111 3947	4217	233963.b.dec g43896	69 4120 451
18 235983.6 dec 2040557	4217	233983.6.dec g43011	16 4129 4500
18 235983.6.dec gessa40	4319 18	200903.0.dec g41119	65 4135 4544
18 235983.6.dec 220574070 3963	4262 18	235983.6.dec g23375	73 4143 4520
10 235983.6.dec 537305014 3370	4503 18	235983.6.dec g380502	21 4145 4500
16 235983.6.dec 05663740 3978	4202 18	235983.6.dec 264121(OT6 4149 4494
18 235983.6.dec 242024414 3978	4224 18	200963.6.dec g821961	4155 4550
18 235983.6.dec 511577714 3978	4197 18	2213930	06 4140 4547
18 235983.6.dec 215221211 3961	4232 18	200903.6.dec g211282	8 4150 4544
18 235983.6.dec 63492500	4210 18	200963.6.dec g298713	8 4150 4540
18 235983.6.dec 634935T0 3982	4533 18	235963.6.dec g497158	0 4156 4500
18 235983.6.dec 215221250 3982	4502 18	200000.0.dec 5070700	
18 235983.6.dec 190300414 3962	4498 18	235963.6.dec g3178678	8 4157 4500
3982	4233 18	235082.0.dec g615559	4157 4541
·		235983.6.dec g615366	4157 4541
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Table 2 cont. 238703.2.dec 1314852H1 238703.2.dec g5526630 238703.2.dec 2508937H1 238703.2.dec g1614392 238703.2.dec 1548794H1 238703.2.dec 2448115T6 238703.2.dec 1734650H1 238703.2.dec 5683758H1 238703.2.d c 1006219H1 238703.2.dec g3754298 238703.2.dec 2042824H1 238703.2.dec 5924675H1 5714657H1 238703.2.dec 238703.2.dec g1761859 238703.2.dec 941045R1 238703.2.dec q831652 238703.2.dec 941045H1 238703.2.dec q3988454 238703.2.dec 225305H1 238703.2.dec g1141926 238703.2.dec 225305R1 238703.2.dec g1615980 238703.2.dec 6269825H1 238703,2.dec q3280478 238703.2.dec 3097424H1 238703.2.dec g3400360 238703.2.dec 602414H1 238703.2.dec g1801195 238703.2.dec 4635313H1 238703.2.dec 2400926H1 238703.2.dec 5597519H1 238703.2.dec g4853169 238703.2.dec 1688380H1 238703.2.dec g2913692 g4900876 238703.2.dec 1688027H1 238703.2.dec 238703.2.dec 4800888H1 238703.2.dec g653486 238703.2.dec 4800701H1 238703.2.dec g518075 238703.2.dec 4321584H1 238703.2.dec g3174834 238703.2.dec 1754493H1 g819357 238703.2.dec 892038H1 238703.2.dec 238703.2.dec g5177747 238703.2.dec 1317383H1 238703.2.dec 889148H1 238703.2.dec 4187641H1 238703.2.dec 2345333H1 238703.2.dec g3181350 238703.2.dec g3240653 238703.2.dec : g2434396 238703.2.dec g3240758 238703.2.dec g1330862 238703.2.dec g2942603 238703.2.dec g4486259 238703.2.dec g1761875 238703.2.dec :: 4663892H1 238703.2.dec 6539614H1 238703.2.dec 6324675H1 238703.2.dec g2056722 238703.2.dec a2216083 238703.2.dec 5271589H1 238703.2.dec q4486257 238703.2.dec q857757 g5637989 238703.2.dec 238703.2.dec q3919555 238703.2.dec q4005327 238703.2.dec g878530 238703.2.dec 4552389H1 238703.2.dec g517836 2046913H1 238703.2.dec 238703.2.dec 5941977H1 238703.2.dec 1380073H1 238703.2.dec g3931604 238703.2.dec g3601039 238703.2.dec 1865544T6 238703.2.dec g3595709 238703.2.dec g2957926 238703.2.dec q2783614 g768855 238703.2.dec 238703.2.dec g3601297 238703.2.dec q690945 238703.2.dec g3840409 238703.2.dec g890047 238703.2.dec g3446295 g5636513 238703.2.dec 238703.2.dec g1721081 238703.2.dec q504660 g2873475 238703.2.dec g1055807 238703.2.dec g884483 238703.2.dec 419939F1 238703.2.dec 238703.2.dec g2675501 238703.2.dec g2932400 238703.2.dec g2464041 238703.2.dec g4649715 238703.2.dec 419939R1 238703.2.dec g760527 238703.2.dec 419939H1 238703.2.dec g2433575 238703.2.dec 5555467H1 238703.2.dec g3840266 g3056119 238703.2.dec g3841172 238703.2.dec 238703.2.dec 1703942H1 238703.2.dec q751598 238703.2.dec g2874018 238703.2.dec 959656H1 238703.2.dec 508789H1 238703.2.dec g919116 3804783H1 238703.2.dec 238703.2.dec 1858793T6 238703.2.dec 3454454H2 238703.2.dec 1573709H1 238703.2.dec 2918331H1 238703.2.dec 6194016H1 238703.2.dec 4416076H1 238703.2.dec 1858793H1 238703.2.dec 5161427H2 238703.2.dec 1858793F6 238703.2.dec 238703.2.dec a696501 g2359502 238703.2.dec g1259095 g5540475 238703.2.dec 238703.2.dec 6380782H1 238703.2.dec g3765011 238703.2.dec 1856905H1 238703.2.dec g2767462 g2004475 238703.2.dec g5127944 238703.2.dec 238703.2.dec g680899 238703.2.dec 941777H1 238703.2.dec q3884338 238703.2.dec 940928T1

18 235983.6.dec 946064H1	Table 2 cont.	j	PCT/US00/25610
235983.6 dec 150640 444	3433 acca		
235983.6 dec 14505505	3436 3640 19 238	703.2.d c g1810435	583 950
10 235983.6 dec 2316000	3452 3849 19 238-	703.2.dec 6166895H	1 500 440
235983.6.dec 145055014	3452 3691 19 339	03.2.dec 4767523H	1 500
235983.6.dec 3661000111	3452 3724 19 2387		1 616 600
235983.6 dec 405750011	040 19 9387		622 896
19 20563.6.dec 6600083H1			628 963
18 335000 6212917H1			628 846
18 20505.B.Gec 5576802H1	3472 2704 19 23870		628 915
18 005005.0.0ec 4649217H1	3478 3740 19 23870	03.2 dec 527000414	630 732
18 00500.0.dec 726102R1	3614 4000 19 23870	3.2 dec 51700 to	630 805
18 00000.0.dec g779370	3485 2000 19 23870	3.2.dec 207400714	650 926
	3486 4004 19 23870	3.2.dec 451000044	651 898
	3485 3000	3.2.dec 220700011	654 914
16 235983.6 dec 2344479	3488 27cc " 200/U	3.2.dec 4460408H1	665 926 665 925
235983.6 dec 220000	3488 2000 13 230/0	3.4.dec 537008H1	
235983 6 dec - 4457	3491 3724 19 33070	P.∠.dec 6385387H1	672 771 676 960
235983.6 dec =20400==	3977 19 239702	7.4.dec 6382968H1	676 905
235983.6 dec 222005	3727 19 239700		681 889
18 335000 4355633H1			702 1006
10 200505.0.dec g877365	15 230700		707 968
18 00500 045305H1	522 265 19 238703	2 dec gossoon	707 ₉₅₂
	545 3700 238703.	2.dec 201320014	709 1074
18 235983.6 dog 20070000 3	547 4447 19 238703	2 100 075150	709 968
10 235983.6 dec 4000400	547 2800 19 238703	2.dec ozenova	713 979
235983.6 dec 12670070	555 2020 19 238/03.2	2.dec 4199245H1	713 983 714 980
235983.6 dec 1367007111 3	38 4072 19 339700	^{2.dec} 1341926F6	740
235983.6 dec appos	3800 19 330703.2	.dec 1338691H1	716 1236 716 996
16 235983.6.dec 01700000	75 3911 19 238702.0	.dec 1341931H1	716 807
18 20383.6.dec 821700H1 36		.uec 1338791H1	716 953
18 205963.6.dec 1908249F6 36	13 2387020		719 921
18 20505.0.dec 3223615H1 35	238703.2	dec 2724040	748 998
19 00500.0.0ec 898523H1 25	98 3974 19 238703.2.	dec 452762714	749 991
18 200000.UEC 898523R1 36/	00 4140 19 238703.2	dec 160004414	768 1014
235983 6 dec 47070	0 3774 19 238703,2,6	dec 3712100111	72 968
10 235983.6 dec 5050640111 30U	2 2000 13 238/03.2	Jec 2544621H1 7	75 1066 85 1027
19 238703.2 dec 1454575111 300	7 2004 '0 208/03.2.0	^{3ec} 857987H1 7	
19 238703.2 dec 611000000 442	695 19 239702.0	iec 4855630H1 p	86 1038 00 1064
19 238703.2.dec 6602392H1 450	13 232700 0		99 1019
19 200703.2.dec 92216250 462	13 222700 6 1		15 1086
10 200703.2.dec g690552 460	726 19 238703.2 d	ec 0010677	5 1082
19 238703.0 4622639H1 469	722 19 238703.2 de	PC 0824400	6 1097
10 95.2.uec 9690544 470	754 ~~~~	C 100646644	
19 238703.2 dec 2651000111 4/0	752 19 238703,2,de	C 4361270114	
19 238703.2 dec 4173000111 469	710 10 238/03.2.de	c g1044402	
2007 W.Z. DPC 0610004		C 2506071H1 83	
19 238703.2 dec 6270000 409	746 19 239703.2.06	C 2041150H1 BAS	.003
19 200705.2.UeC 1564889H1 480	991 19 238702.2.08		1110
19 238703.2.dec 1520074111 409	581 19 238703.2 dec	2070500	1079
2007U3.2.Dec 252050014	238703,2 dec	4972404114	1156
-00703.Z.Dec #1220046	1027 19 238703.2.dec	1242000114	1026
19 200703.2.dec 851725H1 511	774 19 238703.2 dec	4600077111	995
10 222700.2.UEC 852289H1 511	751 19 238703.2.dec	01524407	1122
19 238703.2 dec 15850014 513	820 19 238703,2.dec	260617011	1026
19. 238703.2.dec 870204.D4	785 19 238703.2.dec	169710H1 874	1168
TOO TOO. Z. CIEC TARREST IN	1092 19 238703.2.dec	171492H1 074	1107
= 500700.2.000 1E404001.	831 10 2007U3.2.dec	3244540H1 800	1088 1124
	940 19 238702.0 0	889000R1 800	1456
19 238703.2.dec 3162910H1 560	770 19 238703.0	889000H1 ROO	1158
19 200700.2.dec 3297858H1 573	19 238703.2 dec	2213805H1 891	1111
19 200703.2.dec 1570087H1 594	707 19 238703.2 dec	722196H1 893 g1383346 895	1153
200703.2.dec 1572130H1 ==	79 238703 2 d c	4546400LL	1331
-5.	238703.2.dec	94220214	1162
	75	900 900	1029

				Table	2 cont.	·			
19	238703.2.dec	g2752558	1184	1475	20	038751.5.dec	4649238H1	381	652
19	238703.2.dec	g1735144	1192	1475	20	038751.5.dec	1432680R1	416	909
19	238703.2.dec	1727979H1	1201	1260	20	038751.5.dec	1432680H1	416	681
19	238703.2.dec	1922139H1	1201	1459	20	038751.5.dec	5028237H1	421	526
20 20	038751.5.dec 038751.5.dec	5891158H1 5884165H1	1053 1053	1324 1309	20 20	038751.5.dec 038751.5.dec	3735509H1	428 457	653 759
20	038751.5.dec	g2115213	1053	1513	20	038751.5.dec	g944693 4441762H1	463	75 9 594
20	038751.5.dec	2073762H1	1056	1303	20	038751.5.dec	6599874H1	491	985
20	038751.5.dec	4516520H1	1059	1307	20	038751.5.dec	g1625374	526	843
20	038751.5.dec	2058487H1	1082	1333	20	038751.5.dec	874318R1	530	1138
20	038751.5.dec	g5364468	1107	1573	20	038751.5.dec	874318H1	530	816
20	038751.5.dec	2058968H1	1113	1371	20	038751.5.dec	1692115F6	532	938
20	038751.5.dec	g3665026	1126	1570	20	038751.5.dec	1692115H1	532	749
20	038751.5.dec	g5110105	1129	1563	20	038751.5.dec		540	858
20 20	038751.5.dec 038751.5.dec	5951458H1 1731310T6	1141 1146	1451 1449	20 20	038751.5.dec 038751.5.dec	3838242H1 2701789H1	548 562	828 739
20	038751.5.dec	g4328520	1156	1566	20	038751.5.dec	6615168H1	597	1145
20	038751.5.dec	g657692	1170	1561	20	038751.5.dec	275704H1	600	771
20	038751.5.dec	287003H1	1157	1516	20	038751.5.dec		610	847
20	038751.5.dec	6077974H1	1160	1402	20	038751.5.dec	5562625H1	619	836
20	038751.5.dec	6506745H1	1185	1542	20	038751.5.dec	2061473H1	655	923
20	038751.5.dec	6506945H1	1185		. 20	038751.5.dec	2528543H1	673	973
20	038751.5.dec	3705392H1	1186	1478	20	038751.5.dec	4311336H1	684	1005
20 20	038751.5.dec 038751.5.dec	g3895500 g4329793	1191 1193	1567 1569	20 20	038751.5.dec 038751.5.dec	1731310F6	684 684	1026 908
20	038751.5.dec	g4078804	1199	1570	20	038751.5.dec	1731310H1 4379489H1	695	966
20	038751.5.dec	g2840648	1199	1566	20	038751.5.dec	5317556H1	780	1037
20	038751.5.dec	g2114887	1201	1573	20	038751.5.dec	5316328H1	780	995
20	038751.5.dec	g5368723	1207	1574	20	038751.5.dec	5315861H1	780	931
20	038751.5.dec	g1625271	1207	1566	20	038751.5.dec	5883926H1	847	1111
20	038751.5.dec	g4076938	1213	1571	20	038751.5.dec		847	1103
20	038751.5.dec	g698323	1234	1572	20	038751.5.dec		852	1223
20	038751.5.dec	g2779516	1239	1543	20	038751.5.dec	2348432H1	852	1085
20 20	038751.5.dec 038751.5.dec	1692115T6 g4524126	1244 1246	1529 1570	20 20	038751.5.dec 038751.5.dec	2351121H1 g1512998	852 856	1067 1323
20	038751.5.dec	94453022	1250	1560	20	038751.5.dec		861	1124
20	038751.5.dec	g2212328	1252	1566	20	038751.5.dec	1427102H1	861	1095
20	038751.5.dec	g3896194	1286	1571	20	038751.5.dec	085427H1	874	1061
20	038751.5.dec	g788314	1321	1569	20	038751.5.dec	5177489H1	885	1154
20	038751.5.dec	9723868	1338	1568	20	038751.5.dec	-	910	1134
20	038751.5.dec	g718978	1346	1570	20	038751.5.dec		911	1374
20 20	038751.5.dec	g846280 g1955112	1348 1363	1546	20 20	038751.5.dec	1482576H1	923	1154
20	038751.5.dec 038751.5.dec	g1512999	1365	1566 1580	20	038751.5.dec 038751.5.dec	1005907H1 3700251H1	931 951	1262 1258
20	038751.5.dec	•	1369	1567	20	038751.5.dec		963	1170
20	038751.5.dec	g4617986	1388	1570	20	038751.5.dec	1446007H1	967	1207
20	038751.5.dec	3570081H1	1396	1563	20	038751.5.dec	g1512388	971	1451
20	038751.5.dec	1627963H1	1398	1556	20	038751.5.dec	4674472H1	976	1242
20	038751.5.dec	827450H1	1404	1570	20	038751.5.dec	4677451H1	976	1249
20	038751.5.dec	4467140H1	1441	1566	20	038751.5.dec	2872036H1	976	1204
20 20	038751.5.dec	g3095819 1634593H1	1446	1575	20	038751.5.dec	236814H1	980	1205
20	038751.5.dec 038751.5.dec	1635044H1	1470 1470	1570 1574	20 20	038751.5.dec 038751.5.dec	3986488T6 4215124H1	981 987	1552 1274
20	038751.5.dec	1914206H1	1470	1570	20	038751.5.dec	5085815H1	995	1112
20	038751.5.dec	g4223279	1130	1565	20	038751.5.dec	g751557	996	1232
20	038751.5.dec	g1273271	910	1323	20	038751.5.dec	1470940H1	998	1174
20	038751.5.dec	6064949H1	914	1209	20	038751.5.dec	1634141T6	1019	1524
20	038751.5.dec	6298614H1	1	298	20	038751.5.dec	2073596H1	1021	1241
20	038751.5.dec	4726788H1	82	321	20	038751.5.dec	2351121T6	1020	1526
20	038751.5.dec	667079H1	157	391	20	038751.5.dec	2757071H1	1030	1291
20 20	038751.5.dec 038751.5.dec	3557951H1 9779810	210	476 560	20	038751.5.dec	1634141F6	1026	1571
20	038751.5.dec	g4008516	248 248	568 1570	20 21	038751.5.dec 236099.4.dec	1634141H1 5465756H1	1026 1	1258 288
20	038751.5.dec	2046232H1	307	584	21	236099.4.dec	1622240F6	5	358
20	038751.5.dec	2046232F6	307	607	21	236099.4.dec	1622240H1	5	220
20	038751.5.dec	459067H1	310	561	21	236099.4.dec		· 5	262
20	038751.5.dec	g697893	327	600	21	236099.4.dec	5616026H1	15	· 285
20	038751.5.dec	g711339	359	603	21	236099.4.dec	6171515H1	15	336
				••	78				

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			PCT/US00/25610
19 238703.2.dec 940928H1	Tab	le 2 cont.	1 6 17 6 300 / 256 10
	1335 1470	19 238703 2 dos 2700	
	1349 1461	10 200700.Z.uec 2/62	734H1 45 296
10 2007 00.2.060 93148439	1373 1476	236703.2.dec 2762	735F6 45 144
10 000700.2.080 93740259	1398 1471	236703.2.dec 3082	047H1 45 220
10 200700.2.080 91813211	1415 1477	- 200/US.Z.dec 3337/	00311 50
10 200700.2.000 92/82077	1421 1475	230703.2.dec 2204	257114
10 200.2.000 93891243	1174 1471	-500/UJ.2.dec 3074	745114 50
10 2007 00.2.dec g1801567	1 85	230/03.2.dec 01056	200
238/03.2.dec 4552078H1		19 238703.2 dec 01156	1074
230/U3.2.dec 803067L1		19 238703.2.dec 28500	000
238/03.2.dec 423072Lt		19 238703.2.dec 01616	000
19 238703.2.dec 3148600U4	, , , , , ,	19 238703.2.dec g6536	44 == 001
19 238703.2.dec 353942044		19 238703.2.dec g2007	777 - 270
19 238703.2.dec 358600744		19 238703.2.dec g2056	100
19 238703.2.dec 4434676U4	2 317	19 238703.2.dec 452284	
19 238703.2.dec 327240014	3 277		720
19 238703.2.dec 377377014	5 243		7H1 228 47E
19 238703.2.dec 367225214	5 301	10 200,00.2.000 936951	3 242 656
19 238703.2.dec 40207244	5 199	10 200.2.000 40/429	4H1 261 544
19 238703 2 dog 2004400	6 245	10 200,00.2.080 558512	8H1 261 407
19 238703 2 dec 222500 444	12 542	10 200,00.2.000 434414	8H1 261 470
19 238703 2 dog 24 400	13 276	19 239700.2.060 961380	7 273 546
19 238703 2 dog 2054401	12 294	10 200700.2.080 961382	2 273 540
19 238703.2.dec 265112H1 19 238703.2.dec 4518884H1	11 251	10 000700.2.000 91/618	/4 280 350
	11 227	10 200.2.000 5516366	SH1 296 404
	6 98	10 200.2.080 080407	11 296 540
10 000-50.2.0EC 3449649H1 1	8 282	10 000 Too. 2. uec g1/3524	2 302 500
10 20.2.465 91713059	250	10 1932969	H1 308 556
10 2007 00.2.0ec 4383163H1 2	5 163	10 000-00.2.000 0485648	H1 316 945
		236703.2.dec 1932969	F6 308 700
10 3482476H1 2	6 298	238703.2.dec 5810554	H1 309 636
10 2007.2.000 1865544F6 2		238/03.2.dec 5810747	H1 309 624
10 100.2.dec 30/4630H1 2/	300	430/03.2.dec 5150576	U4 04=
10 000-100-2-UEC 1865544H1 24	5 276	230/03.2.dec 035/63u	4 111 0/1
10 236703.2.dec g1722043 21		13 438/03.2.dec 030530H	000
236703.2.dec 482687H1 26		238/03.2.dec 63131371	14 023
10 2007/US.2.dec 3747723H1 26		19 238703.2.dec g884522	2007
238703.2.dec 4569176H1 20		19 238703.2.dec 033396U	
236703.2.dec 486383H1 27		19 238703.2.dec 033287H	700
10 236703.2.dec 1338949H1 28	270	19 238703,2,dec 073634LI	040
10 236703.2.dec 3216994H1 33	257	19 238703.2.dec 073602H1	- 040
236703.2.dec 2935963H1 31	118	19 238703.2.dec 3282162H	4
236703.2.dec 485050H1 33	270	19 238703.2.dec 4398152L	4
230/03.2.dec 5165752U4 00		19 238703.2.dec 0616510	
10 236703.2.dec 484526H1 33	280	19 238703.2.dec 183426411	378 587
230/03.2.dec 486726U4	266 266	19 238703.2.dec 4917182H	, 510 001
230/03.2.dec 4152420U4 -	313	19 238703.2.dec 187622L1	
236703.2.dec 3592529H1 36		19 238703.2.dec 24780774	386 569
238703.2.dec 3174680H1 36	347	19 238703.2.dec 2008040H	1001
238703.2.dec 799668H1 35	291 260	19 238703.2.dec 2488430H	003
10 236703.2.dec 3614882H1 36	269	19 238703.2.dec 6064576LI4	
230/03.2.dec 28000ecu4 0-	329	19 238703.2.dec 5905782U4	
19 238703.2.dec 4802649H1 07		19 238703.2.dec 01958061	
19 238703.2.dec 2800052H1 00		19 238703.2.dec g890104	412 842
230/03.2.dec 01750652 04		19 238703.2.dec 4596304Lt	412 540
238/03.2.dec 4010722H4 02	381	19 238703.2.dec 4597956H1	412 665
230/03.2.dec 24/55/50/		19 238703.2.dec go18642	412 662
- 10 230/03.2.dec 346200204	292		421 642
19 238703.2.dec 4911150U4	186	000000	421 635
200/03.2.dec 3358571U4	338		423 668
19 238703.2.dec 244811556	229 1	0 200700.2.000 9104632/	426 781
19 238703.2.dec 2449115U4	391 1		434 735
19 238703.2.dec 3538100H1	290 1		441 744
19 238703.2.dec 3504412U4	338 ₁	000700.2.dec 4900130H1	441 704
19 238703.2.dec d833330	353 1		442 913
19 238703.2 dec 01761950	472		1207 1449
19 238703.2.dec 3151500U4 42	380 1	3548126	1179 1477
19 238703 2 dec 4075000111	326	9861832	1177 1470
19 238703 2 d c 237375711 45	322	0098//H1	1181 1411
19 238/03.2.d c 3273757H1 45	310 19	2207020	1184 1486
	- '-	238703.2.d c g1750539	1181 1478
	77		

	Table 2 cont.								
21	236099.4.dec	6095067H1	19	323	21	236099.4.d c	3251609F6	1097	1514
21	236099.4.dec	4634203H1	19	302	21	236099.4.dec	3251609H1	1097	1401
21	236099.4.dec		21	271	21	236099.4.dec	5865657H1	1100	1367
21	236099.4.dec	g2080681	19	376	21	236099.4.dec	2512120T6	1105	1670
21	236099.4.dec	4415334H1	23	287	21	236099.4.dec	5744630H1	1113	1381
21	236099.4.dec		24	207	21	236099.4.dec	g2161810	1118	1554
21 21	236099.4.dec	6549766H1	25	620	21	236099.4.dec	1600152H1	1137	1343
21	236099.4.dec 236099.4.dec	3566591H1 1389095H1	25 25	264	21	236099.4.dec	5605484H1	1156	1430
21	236099.4.dec	6008955H1	25 27	250 309	21 21	236099.4.dec	g2027235	1171	1452
21	236099.4.dec	2070696H1	27	280	21	236099.4.dec 236099.4.dec	5314568H1 g4260562	1213 1231	1433 1706
21	236099.4.dec	3746033H1	30	283	21	236099.4.dec	4180560T8	1231	1685
21	236099.4.dec	3586436H1	29	222	21	236099.4.dec	g2539107	1236	1709
21	236099.4.dec	3750032H1	30	303	21	236099.4.dec	3805461H1	1240	1547
21	236099.4.dec	2818544H1	33	327	21	236099.4.dec	g2566268	1240	1708
21	236099.4.dec	4762420H1	33	291	21	236099.4.dec	g4982919	1239	1703
21	236099.4.dec	2655186H1	37	328	21	236099.4.dec	2411813H1	1243	1484
21 21	236099.4.dec 236099.4.dec	258736H1	39	131	21	236099.4.dec	5302238H2	1247	1500
21	236099.4.dec	945328H1 238507R1	42 45	312 535	21	236099.4.dec	g3417716	1248	1706
21	236099.4.dec	3219002H1	44	336	21 21	236099.4.dec 236099.4.dec	g4311838 g4085716	1248	1710
21	236099.4.dec	3808753H1	47	351	21	236099.4.dec	1926463R6	1248 1249	1628 1595
21	236099.4.dec	1388805H1	47	293	21	236099.4.dec	6375524H1	1249	1545
21	236099.4.dec	5421931H1	48	300	21	236099.4.dec	g1404330	1248	1691
21	236099.4.dec	238507H1	46	275	21	236099.4.dec	1926463T6	1249	1661
21	236099.4.dec	1003290H1	48	274	21	236099.4.dec	1802927T6	1250	1657
21	236099.4.dec	2698971H1	48	261	21	236099.4.dec	1926463H1	1249	1463
21	236099.4.dec	3148652H1	48	338	21	236099.4.dec	g2194686	1251	1505
21 21	236099.4.dec 236099.4.dec	2893513H1 5095847H1	51 54	325 320	21	236099.4.dec	g2194934	1251	1444
21	236099.4.dec	6125638H1	53	520 536	21 21	236099.4.dec 236099.4.dec	4058927H1	1256	1530
21	236099.4.dec	2457030H1	53	283	21	236099.4.dec	g5636939 772821H1	1256 1263	1706 1474
21	236099.4.dec	3535276H1	54	344	21	236099.4.dec	238507F1	1272	1699
21	236099.4.dec	3590255H1	54	320	21	236099.4.dec	2239910H1	1275	1483
21	236099.4.dec	6119241H1	65	406	21	236099.4.dec	2603359T6	1280	1669
21	236099.4.dec	6125460H1	65	464	21	236099.4.dec	g2080682	1286	1711
21	236099.4.dec	6119192H1	65	626	21	236099.4.dec	4602424T6	1289	1665
21 21	236099.4.dec 236099.4.dec	4194343H1	73	374	21	236099.4.dec	6106532H1	1292	1631
21	236099.4.dec	4158894H1 g389440	94 144	350 513	21	236099.4.dec	g4261108	1296	1706
21	236099.4.dec	g570803	175	522	21 21	236099.4.dec 236099.4.dec	g4223524 g4109832	1298	1706.
21	236099.4.dec	g673306	175	488	21	236099.4.dec	1622240T6	1300 1301	1706 1667
21	236099.4.dec	4908387H1	180	442	21	236099.4.dec	g2161667	1311	1709
21	236099.4.dec	641175H1	225	469	21	236099.4.dec	g3432780	1309	1706
21	236099.4.dec	3790670H1	278	569	21	236099.4.dec	g2350629	1312	1706
21	236099.4.dec	5268416H1	316	569	21	236099.4.dec	g3898751	1316	1706
21	236099.4.dec	3589959H1	375	703	21	236099.4.dec	•	1319	1699
21 21	236099.4.dec 236099.4.dec	4190221H1	397	676 601	21	236099.4.dec	g5177660	1319	1696
21	236099.4.dec	3967058H1 1427256F6	408 502	691 1075	21 21	236099.4.dec 236099.4.dec	4842322H1	1321	1602
21	236099.4.dec	1427256H1	502	737	21	236099.4.dec	J - · · ·	1322 1324	1699 1599
21	236099.4.dec	5386176H1	528	657	21	236099.4.dec		1347	1712
21 .	236099.4.dec	1718014H1	655	897	21	236099.4.dec	4402183H1	1356	1611
21	236099.4.dec	4010711H1	678	949	21	236099.4.dec	g5674364	1357	1707
21	236099.4.dec	6542821H1	777	1331	. 21	236099.4.dec	g4153286	1360	1699
21	236099.4.dec	5492621H1	894	1166	21	236099.4.dec		1368	1626
21	236099.4.dec	5492721H1	894	1164	21	236099.4.dec	•	1381	1699
21 21	236099.4.dec	6256961H1	910	1160	21	236099.4.dec		1391	1706
21	236099.4.dec 236099.4.dec	5979837H1 5273408H1	946 983	1230	21	236099.4.dec		1401	1707
21	236099.4.dec	3163001H1	994	1239 1276	21 21	236099.4.dec 236099.4.dec		1403	1705
21	236099.4.dec	g2032366	994	1273	21	236099.4.dec	501997H1 6350508H2	1406	1617
21	236099.4.dec	6308691H1	1006	1558	21	236099.4.d c	6366728H1	1409 1415	1706 1682
21	236099.4.dec	g2743130	1035	1393	21	236099.4.d c	g2715316	1418	1706
21	236099.4.dec	5554939H1	1061	1304	21	236099.4.d c		1433	1661
21	236099.4.dec	1427256T6	1088	1660	21	236099.4.dec	2365114H1	1433	1660
21	236099.4.dec	5224984H1	1093	1335	21	236099.4.dec		1439	1699
21	236099.4.dec	5560308H1	1096	1332	21	236099.4.dec	3168773H1	1441	1705
					79				

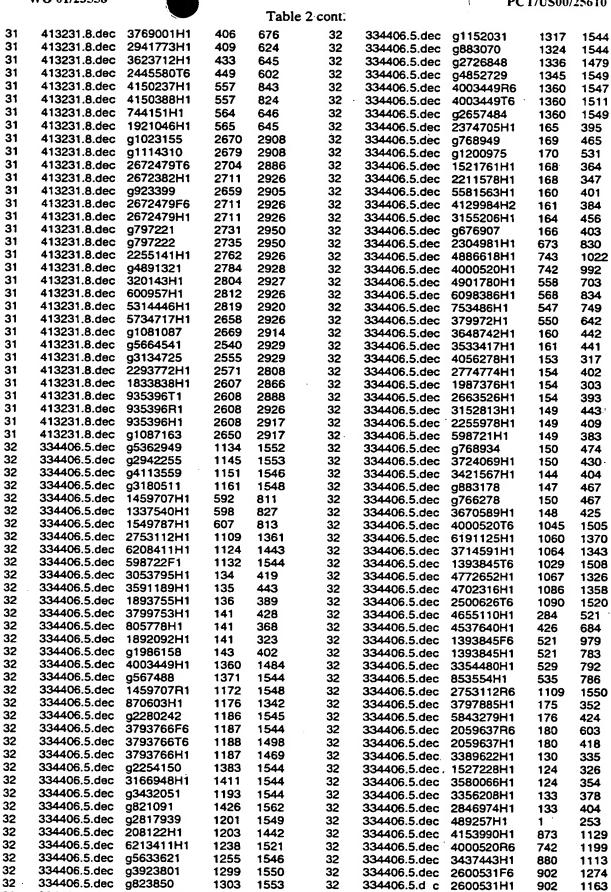
Table 2 cont. Table	W O 01/23558			
2 2 30099 4.0e 0 1982078 1 1482 1850 2 4 466521.6.de 450289H1 1 249 212 220094.4ce 0 1982078 1 1488 1650 2 4 466521.6.de 2 343657F6 3 202 212 230099 4.0e 0 1982078 1 1503 1983 2 4 466521.6.de 2 3465296 8 322 212 236099 4.0e 0 1982099 1 1505 1706 2 4 466521.6.de 2 3465978 2 2 2 2 35099 4.de 0 1982099 1 1505 1706 2 4 466521.6.de 3 365049H1 9 209 2 2 2 35099 4.de 0 1982099 1 1505 1706 2 4 466521.6.de 3 365049H1 9 209 2 2 2 35099 4.de 0 1982099 1 1508 1706 2 4 466521.6.de 3 365049H1 9 209 2 2 3 2 3 2 3 3 3 3 3 3 3 3 3 3 3 3 3		Та	ble 2	PCT/US00/25610
21 236099 4.dec			. /	2.010
22 22 23 23 24 24 25 25 25 25 25 25 25 25 25 25 25 25 25	21 236099.4.dec 01626759		+00521.6.dec	455020014
2 250099 4.de	21 235099.4.dec 2750215114		²⁴ 466521.6.dec	6141664114
22 236099 4.dec 9342605 1498 1593 924 4665216.dec 9092590H1 8 200 21 236099 4.dec 9705688 1503 1939 24 4665216.dec 4092590H1 9 200 21 236099 4.dec 94201811 1503 1508 1706 24 4665216.dec 9436479H2 10 276 276 276 276 276 276 276 276 276 276	230099.4.d c 3950250H1	1000	²⁴ 466521.6.dec	234505750 0 205
221 236099 4.dec 9199999 1508 1706 24 466521.6.dec 3636049H1 9 270 212 236099 4.dec 9199999 1508 1706 24 466521.6.dec 3636049H1 9 270 243 243 243 243 243 243 243 243 243 243	236099.4.dec g3842605	1000	²⁴ 466521.6.dec	01068000 - TUS
2 2 23099 4.dec 9 399989 1508 1706 24 466521.6.dec 485147942 10 278 279 279 279 279 279 279 279 279 279 279	23 236099.4.dec g2705688	1000	²⁴ 466521.6.dec	409250004
221 220093 4.dec 942018H1 1513 1706 24 466521.6.dec 84951479H2 10 249 221 23093 4.dec 942018H1 1513 1706 24 466521.6.dec 922794H1 12 230 241 230094.dec 942018H1 1513 1706 24 466521.6.dec 922794H1 12 230 241 230094.dec 942018H1 1513 1706 24 466521.6.dec 922794H1 12 230 241 230094.dec 942018H1 1513 1706 24 466521.6.dec 922794H1 12 230 241 230094.dec 942018H1 1513 1706 24 466521.6.dec 922794H1 12 230 241 230094.dec 942018H1 1513 1706 24 466521.6.dec 92008042 20 339 261 21 2360994.dec 9456764 1559 1705 24 466521.6.dec 92008042 20 339 261 21 2360994.dec 9456764 1559 1930 24 466521.6.dec 92008042 29 339 21 2360994.dec 9456764 1559 1930 24 466521.6.dec 945664 13368H1 1688 1706 24 466521.6.dec 945664 173368H1 1688 1706 24 466521.6.dec 945664 173669 139144 1627 1932 24 466521.6.dec 945662 1369 139144 1627 1932 24 466521.6.dec 945662 1369 139144 1627 1932 24 466521.6.dec 945662 1369 139144 1627 1932 24 466521.6.dec 94569 139144 1627 1932 24 466521.6.dec 94569 139144 1627 1932 24 466521.6.dec 94569 13914 1932 1935 24 466521.6.dec 94569 13914 1932 1932 24 466521.6.dec 94569 13914 1932 1935 1932 1932 1932 1932 1932 1932 1932 1932	21 236099.4.dec g3191411		²⁴ 466521.6.dec	3636040114
221 23099 4.0ec 942018T1 1513 1706 24 466521.6.dec 822739H1 10 2329 221 232999 4.0ec 942018T1 1513 1706 24 466521.6.dec 3800786H1 12 2329 221 232999 4.0ec 476498271 1513 1706 24 466521.6.dec 4399258H1 1519 1725 24 466521.6.dec 4399258H1 152 2329 221 232999 4.0ec 942018T1 1519 1725 24 466521.6.dec 4399258H1 19 284 221 232999 4.0ec 94587742 1532 1699 24 466521.6.dec 4399258H1 19 284 221 232999 4.0ec 94587742 1532 1699 24 466521.6.dec 439928H1 19 284 221 236099 4.0ec 94587742 1551 1709 24 466521.6.dec 439928H1 19 284 221 236099 4.0ec 173758H1 1585 1857 24 466521.6.dec 439928H1 19 284 284928H1 19 284928H1 19 284928H1 19 284 284928H1 19 284928H1 19 28	23 236099.4.dec g1990989		²⁴ 466521.6.dec	1951/7040
221 230099 4.0e 942018F1 1513 1688 24 466521.6.de 322794H1 12 329 221 230099 4.0e 6409055H1 1513 1708 24 466521.6.de 323465H1 16 266 224 221 230099 4.0e 6409055H1 1519 1725 24 466521.6.de 329928H1 16 266 221 230099 4.0e 92539580 1535 1699 24 466521.6.de 3290804 20 329 221 230099 4.0e 92539580 1535 1699 24 466521.6.de 3290804 10 22 32 32 32 32 32 32 32 32 32 32 32 32	235099,4,dec 042010U4	4	²⁴ 466521.6.dec !	584364004
23 230099 4.0ec 942018F1 1519 1706 24 466521.6.dec 329268611 19 289 24 466521.6.dec 329268611 19 289 24 466521.6.dec 32926861 19 289 24 466521.6.dec 329268611 19 289 24 466521.6.dec 32926861 19 389 39 39 39 39 39 39 39 39 39 39 39 39 39	236099.4.dec 942018T1	4.5.	24 466521.6.dec 9	22204114
2.2	23 236099.4.dec 942018R1	4 =	²⁴ 466521.6.dec 3	RRATORUA
23 230099.4.dec 2377067F1 1531 1711 24 465521.6.dec 23008042 20 230	236099.4.dec 6409055H1	4	24 466521.6.dec 6	25AGGGUA
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21 236094 dec 2736471 30 2312 236094 dec 2736471 30 2312 236094 dec 2736471 30 2312 236094 dec 27376471 48 216 236094 dec 27376471 48 216 236094 dec 273766871 48 271 236094 dec 273766871 48 271 236094 dec 273766871 48 271 236094 dec 27366871 48 271 236094 dec 2736094 de	230099.4.dec 4784982H2		²⁴ 466521.6.dec g	2008042 20 200
21 236094 d. dg	230099.4.dec g2539580	4	²⁴ 466521.6.dec 4	746004110
21 226093.4 dec 2151758911 1585 1867 24 466521.6.dec 15101H1 48 307 2210293094.dec 21315891 1585 1867 24 466521.6.dec 15101H1 48 307 236093.4 dec 1733669H1 1588 1706 24 466521.6.dec 15101H1 49 279 21 236093.4 dec 1733669H1 1602 1992 24 466521.6.dec 234505776 295 21 236093.4 dec 1733669H1 1602 1992 24 466521.6.dec 234505776 295 21 236093.4 dec 1733669H1 1602 1992 24 466521.6.dec 234505776 295 21 236093.4 dec 1733669H1 1602 1992 24 466521.6.dec 234505776 295 21 236093.4 dec 1733669H1 1634 1699 24 466521.6.dec 2364518-H1 491 769 21 236093.4 dec 1733669H1 1701 1883 24 466521.6.dec 2364518-H1 491 769 21 236093.4 dec 1870.4 d	24 220033.4.uec 92/82664		** ***********************************	3726404 00
21 236093.4 dec 1733669H1 1588 1706 24 465521.6.dec 4976429H1 51 249 249 249 249 249 249 249 249 249 249	24 94567742	4.00	400521.6.dec 3	439046H1 38 316
21 236094.dec 735668H 1588 1706	24 2151/58H1		24 400521.6.dec 30	359046H1 48 307
21 236099.4.dec 92816105 1602 1992 24 466521.6.dec 234505776 295 796 21 236099.4.dec 92816105 1602 1992 24 466521.6.dec 234505776 295 796 21 236099.4.dec 95028373H1 1634 1699 24 466521.6.dec 2356970H1 451 736 21 236099.4.dec 9562487 1682 1983 24 466521.6.dec 3358415H1 497 21 236099.4.dec 9562487 1682 1983 24 466521.6.dec 5549567H1 495 792 21 236099.4.dec 9570491 1701 1983 24 466521.6.dec 5549567H1 495 792 22 350875.2.dec 23342386 1 2011 24 466521.6.dec 245878HH1 240 24 24 24 24 24 24 2	24 2000.4.dec 1/33668H1		400521.6.dec 10	05101H1 49 240
21 236093.4.dec 21 236093.4.de	24 1/33668F6		24 400521.6.dec 49	76429H1 51 300
21 226093.4.dec 50228373H1 1634 1699 24 465521.6.dec 33789416H 447 799 778 236093.4.dec 2306752.2.dec 236093.4.dec	24 92816105		24 400521.6.dec 23	45057T6 205 700
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21 236099.4.dec 9552487 1682 1983 24 466521.6.dec 5549503H1 491 764 912 1236098.4.dec 9552487 1682 1983 24 466521.6.dec 5549503H1 493 739 21 236098.4.dec 9552487 1682 1983 24 466521.6.dec 5549503H1 495 732 2350875.2.dec 3350875.2.dec 33308230H1 1 201 24 466521.6.dec 1208881R1 649 834 350875.2.dec 3350875.2.dec 344204H1 8 308 24 466521.6.dec 1208881R1 649 834 350875.2.dec 2324423H6 15 473 24 466521.6.dec 2535788H1 649 834 350875.2.dec 393129H1 19 366 24 466521.6.dec 244602H1 690 834 22 350875.2.dec 939129H1 19 366 24 466521.6.dec 1208881R1 649 834 24 466521.6.dec 3350875.2.dec 939129H1 19 190 465 24 466521.6.dec 5027483H1 704 834 22 350875.2.dec 339129H1 19 190 465 24 466521.6.dec 5027483H1 704 834 22 350875.2.dec 1387573H1 22 334 24 466521.6.dec 5027483H1 704 834 22 350875.2.dec 1387573H1 22 334 24 466521.6.dec 5027483H1 704 834 22 350875.2.dec 1387573H1 22 344 24 466521.6.dec 5027483H1 704 834 22 350875.2.dec 1388319H1 22 150 24 466521.6.dec 5027483H1 704 834 22 350875.2.dec 1388319H1 22 150 24 466521.6.dec 3729373H1 704 834 22 350875.2.dec 1388319H1 22 150 24 466521.6.dec 3729291T1 732 792 23 350875.2.dec 94622245 54 467 24 466521.6.dec 3729291T1 732 792 23 350875.2.dec 94622245 54 467 24 466521.6.dec 3729291T1 732 792 23 350875.2.dec 94622245 54 467 24 466521.6.dec 3729291T1 732 792 23 350875.2.dec 94622245 54 467 24 466521.6.dec 3729291T1 732 792 23 350875.2.dec 94622245 54 467 24 466521.6.dec 3729291T1 732 792 23 350875.2.dec 94022245 54 467 24 466521.6.dec 3729291T1 732 792 350875.2.dec 94022245 54 467 24 466521.6.dec 3729291T1 732 792 350875.2.dec 94022245 54 467 24 466521.6.dec 3729291T1 732 792 350875.2.dec 94022245 54 467 24 466521.6.dec 3729291T1 732 792 350875.2.dec 94022245 54 467 24 466521.6.dec 3729291T1 732 792 350875.2.dec 94022245 54 467 24 466521.6.dec 3729291T1 732 792 350875.2.dec 94022245 54 467 24 466521.6.dec 3729291T1 732 792 350875.2.dec 94022245 54 467 24 466521.6.dec 3729291T1 732 792 350875.2.dec 94022245 54 467 24 466521.6.dec 3729374H1 748 324 466521.6.dec 3729374H1 748 324 466521	24 = 5028373H1		400521.6.dec 48	39970H1 451 736
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23 466521.5.dec g2955287 388 712 25 474522.8.dec 3108601H1 693 863 24 466521.6.dec 2345057H1 3 257 25 474522.8.dec 6060467H1 700 1291 24 466521.6.dec 4788225H1 1 213 25 474522.8.dec 6264353H1 708 1290 25 474522.8.dec 4246633H1 708 1290	23 466521.5.dec g1395834 382	740	25 474522.8.dec 428747	N. 14
24 466521.6.dec 4788225H1 1 213 25 474522.8.dec 6060467H1 700 1291 213 25 474522.8.dec 4246633H1 708 1290	23 466521.5.dec g2955287 389	710	**************************************	Ille oo-
213 25 474522.8.dec 6264353H1 708 1290	24 400521.6.dec 2345057H1 3	0.00	5 474522.8.dec 606046-	7114 700
25 474522.8.dec 4246633H1 707	400521.6.dec 4788225H1 1	040	" 4/4022.8.dec 6264252	olid man
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25	474522.8.dec	4540246H1	733	907	25	474522.8.dec	4934454H1	74	040
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25	474522.8.dec					474522.8.dec	3297190H1	105	348
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25	474522.8.dec	6132989H1	778	992	25	474522.8.dec	g3415484	1	284
25	474522.8.dec	g1509952	781	970	25	474522.8.dec	g4188698	1	253
25	474522.8.dec	1358904T6	829	1408	25	474522.8.dec	g2003036	1	375
25	474522.8.dec	1643034F6	837	1404	25	474522.8.dec	g3923493	1	413
25	474522.8.dec	1643034H1	837	1062	25	474522.8.dec	3741633H1	. 1	285
25	474522.8.dec	3013573H1	847	1131	25	474522.8.dec	6397326H1	9	300
25	474522.8.dec	3683622H1	851	1148	25	474522.8.dec	g5395341	12	454
25	474522.8.dec	1623584H1	849	1068	25	474522.8.dec	2497236F6	16	475
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25	474522.8.dec	6304964H1	1033	1418	26	231583.3.dec	g2742647	1165	1261
25	474522.8.dec	1227119H1	1052	1287	26	231583.3.dec	2605033H1	1170	1261
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25	474522.8.dec	3659596H1	1143	1365	26	231583.3.dec	g1874192	86	540
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25	474522.8.dec	5710431H2	189	398			-		586
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25	474522.8.dec	550275H1	255	477	26	231583.3.dec	g3308494	819	1256
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25	474522.8.dec	4545853H1	402	642	26	231583.3.dec	3452837H1	849	899
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25	474522.8.dec	2277358H1	406	689	26	231583.3.dec	g4988189	867	1264
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25	474522.8.dec	2497236H1	16	330	26	231583.3.dec	3794634H1	136	418
25	474522.8.dec	5862690H1	23	284	26	231583.3.dec	1420279H1	171	439
25	474522.8.dec	g3785848	60	276	26	231583.3.dec	2967782F6	97	607
25	474522.8.dec	g3144117	60	434	26	231583.3.dec	3296842H1	98	362
25	474522.8.dec	g2809679	60	412					
25	474522.8.dec	g1691199	60		26 26	231583.3.dec	4551367H1	108	354
25 25	474522.8.dec			409	26 26	231583.3.dec	2608335F6	108	417
25		g2183371	60	395	26 26	231583.3.d c	2608335H1	108	358
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25	474522.8.dec	g795882	59	322	26	231583.3.dec	3585168H1	117	304
25	474522.8.dec	g3181844	60	244	26	231583.3.dec	g1637120	114	455
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25	474522.8.dec	g2880771	66	363	. 26	231583.3.dec	g4003920	783	1181

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26 231583.3.dec 2644845H1		able 2 cont. PC 1/US00/25610
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26	497 629	27 213051.5.dec 1909813F6 982 1566
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26 25.5.050 91558633	538 923	27 2.5051.5.dec 4541904H1 986 1251
26 0015-0-0-0-0-0-0-0-0-0-0-0-0-0-0-0-0-0-0	591 735	27 215051.5.dec 5154081H1 1013 1365
26 3/184//H1	621 925	27 213051.5.dec 5951978H1 1040 1254
20 1740/14R6	672 1076	27 215051.5.dec 5947813H1 1040 1300
26 -01000.3.dec 1/40/14H1	672 900	27 213051.5.dec g1757881 1045 1209
26	693 945	27 215051.5.dec 4948780H1 1054 1329
3/99139H1	694 985	27 215051.5.dec 1704449H1 1069 1277
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26 296//8216	733 1356	27 213051.5.dec 1855956F6 1105 1660
20 == 000.0.dec 299305/H1	755 1011	27 215051.5.dec 1855956H1 1105 1270
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26 231583.3.dec g1897911 26 231583.3.dec g2570924	87 535	27 215051.5.dec 6483737H1 1145 1660
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3206034H1	61 312	27 213051.5.dec 6348133H1 1166 1290
27 215051 5 de 000083M7	1576 1827	27 215051.5.dec 5048814H1 1250 1450
27 04501.5.060 93330058	1585 1975	27 213051.5.dec g4891899 1271 1672
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07 11.0.dec 91/301H1	1592 1889	27 213031.5.dec 4710136H1 1286 1272
37 94435/60	592 1971	27 215051.5.dec 4198162H1 1288 1514
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27 215051.5 doc = 109036216 1	604 1934	213051.5.dec 1868512H1 1301 1575
27 22001509 1	608 1971	27 213031.5.0eC 1868318H1 1301 1562
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27 215051 5 dog 3010494 10	647 1971	27 215051.5.dec 853710H1 1325 1519
27 215051.5 dec 121000000	709 1936	27 215051.5.dec 1855956T6 1332 1036
27 215051 5 dog 50000 to 17	709 1953	27 215051.5.dec 4699841T6 1338 1945
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2/ 215051,5,dec 04125100 47		27 215051 5 dec 1210132H1 1411 1546
2/ 215051 5 dec 44044		2/ 215051 5 dog 200005000 1472 104/
2/ 215051.5.dec 5710048H1 404	_	27 215051.5.dec 189055050 1419 1933
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2/ <15051.5.dec g2805404		27 215051.5 dog 500705011 1440 1930
2/ 215051.5.dec 352110344 705	• • •	E'
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27 275051.5.dec 1301667Ee 000	1113	27 215051 5 dog 5703457 1503 1759
2/ 2/3031.5.dec 1301667L1 000	1216	27 215051 5 dog 570015H1 1504 1973
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27	215051.5.dec	g5112194	1520	1979	28	277726.5.dec	3349217H1	168	338
27	215051.5.dec	5205618H1	1529	1639	28	277726.5.dec		169	427
27	215051.5.dec	g2705032	1533	1973	28	277726.5.dec		174	461
27	215051.5.dec		1536	1971	28	277726.5.dec		174	454
27	215051.5.dec		1537	1782	28	277726.5.dec		186	396
27	215051.5.dec	g2805413	1541	1971	28	277726.5.dec	3506561H1	190	504
27	215051.5.dec	•	1548	1973	28	277726.5.dec		190	391
27	215051.5.dec	-	1566	1971	28	277726.5.dec		191	320
27	215051.5.dec	3218687H1	1	269	28	277726.5.dec	6550846H1	211	793
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27 27	215051.5.dec	686080H1	5	246	28	277726.5.dec		272	543
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27	215051.5.dec	3333490H1	33	291	28 28	277726.5.dec	1856677H1	305	577
27	215051.5.dec	1554835H1	54	250	28 28	277726.5.dec 277726.5.dec		391	640
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27	215051.5.dec	3333365H1	55	317	28	277726.5.dec	3138453H1	488	731 778
27	215051.5.dec	4346748H1	139	305	28	277726.5.dec	2941152H1	493	773
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27	215051.5.dec	4103113H1	275	556	28	277726.5.dec	623556H1	586	826
27	215051.5.dec	3529273H1	387	659	28	277726.5.dec	4902769H1	626	899
27	215051.5.dec	1428015H1	475	675	28	277726.5.dec	2265514H1	653	897
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28	277726.5.dec	g776275	126	396	28	277726.5.dec	1493174H1	833	1070
28	277726.5.dec		132	665	28	277726.5.dec	3520394H1 2076038H1	843 868	1093
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28	277726.5.dec	6134663H1	145	414	28	277726.5.dec	4904840F6	956	1187
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28	277726.5.d c	1996357H1	167	442	28 29	277726.5.dec 978637.1.dec	4338458H1	1132	1228
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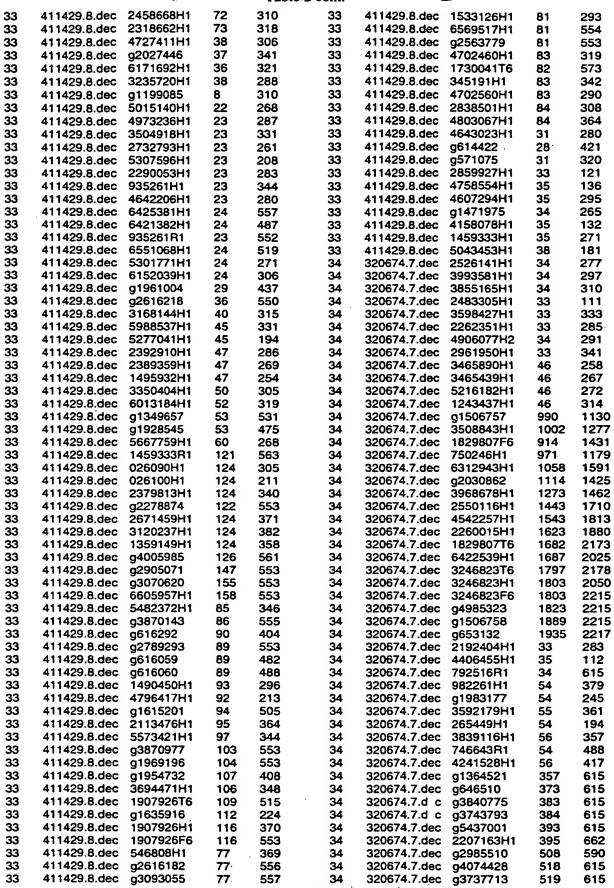
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31 413231.8.dec 6538992H1 1965 2556 31 413231.8.dec 1321333F6 1076 1424 31 413231.8.dec 91024057 1997 2356 31 413231.8.dec 3039505H1 1083 1346 31 413231.8.dec 92006123 2041 2484 31 413231.8.dec 91149424 1090 1424 31 413231.8.dec 97803H1 2122 2345 31 413231.8.dec 6597349H1 110 637 31 413231.8.dec 1613405H1 2122 2345 31 413231.8.dec 3615019H1 327 624 31 413231.8.dec 1613405H1 2122 2324 31 413231.8.dec 5101007H1 328 572 31 413231.8.dec 5330251H1 2158 2405 31 413231.8.dec 2728285H1 335 582 31 413231.8.dec 5330251H1 2158 2427 31 413231.8.dec 983374H1 355 641 31 413231.8.dec 2154518H1 226 2510	31 4	13231.8.dec 9	90688H1 102			31 47	3231.8.dec 13	2404014	72 1426
31 413231.8.dec g2006123 2041 2484 31 413231.8.dec g1149424 1090 1	31 4	13231.8.dec 6	38992H1 196		^	31 41	3231.8.dec 13	24222	
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31 413231.8.dec 5330251H1 2158 2427 31 413231.8.dec 983374H1 355 641 31 413231.8.dec 2285744H1 2218 2429 31 413231.8.dec 6368807H1 362 627 31 413231.8.dec 2154518H1 2226 2510 31 413231.8.dec 2271165H1 2305 2571 31 413231.8.dec 2382180H1 373 637 31 413231.8.dec 26685512H1 2323 2606 31 413231.8.dec 2061273H1 378 638	31 47	3231.8.dec 534	20447114	2561			231.8.dec 510	1007H1 328	
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32 334406.5.dec 2600531T6	12	ble 2 cont.	1 € 17 € 300/25610
	920 1504	33 411420 9 40-	
20 100.0.0ec 103//93HT	939 1147	, 411429.8.dec ose	30906 250 562
30 30,700,3,060 3934/30H1	760 1046	711469.8.000 627	33040
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32 334406.5.dec 199393976		31 411420 0	700.40
32 334406.5.dec 1993930FC	961 1506		76646 268 554
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20 1993838H1	975 1179	22 444	1032 205 550
20 000.0.dec 000/592H1	986 1515	22 444 25.0.066 922	36485 301 507
30 100Z119H1	1009 1189	411429.8.dec n310	11070
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32 334406.5.dec 5016037U4	4		
32 334406.5.dec 332846044	190 411		5125 319 556
32 334406.5.dec 879416H1	190 468		42H1 323 562
	160 388	22 11.723.0.000 0555	57H1 324 510
	160 396	22 11723.0.086 2338	85H1 327 552
20 2230254H1	160 418	₩ 411429.8.dec a307/	2610
20 00 1100.5.dec 2//4782H1	154 394	411429.8.dec 14054	SOCIA SOCIA
32 334406.5.dec 3271675H1	4	33 411429.8.dec 01300	1675
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32 334406.5.dec 250062656	122 339		202 388 764
32 334406.5.dec 152704044	122 551		07H1 391 553
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	1294 1592	33 711429.8.dec g1101	419 396 622
22 210/23376	1445 1989	411429.8.dec 0570E	00
20 11-23.0.Uec 8/1885H1	1449 1604	411429.8.dec 03434	866
20 11-23.6.0ec 210/233R6	1452 1942	33 411429.8.dec 04283	207
411429.8.dec 2107222U4	4.400	33 411429.8.dec 2080ec	1114
33 411429.8.dec 4935576H4	445		
33 411429.8.dec 2295977H1	1154 1419	33 411429.8.dec 293320 411429.8.dec 459243	05H2 469 527
33 411429.8.dec 0565005	1239 1493		18H1 488 557
	1538 1879	22 11425.0.060 186033	10F6 498 074
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32 A44 120.0.0ec 3/2/538H1	1759 2035	22 411429.8.dec 839111	H1 519 577
22 11 20.0.000 01952/3H1	1828 2124	411429.8.dec 451756	2014
23 711429.8.dec 4981402H1	2013 2256	33 411429.8.dec 459076	L1 070
411429.8.dec 4981202U4		33 411429.8.dec 443671	ELI4 ATT
411449.8.dec 2721111111		33 411429.8.dec 052139	
33 411429.8.dec 3216954114			
33 411429.8.dec 01774607	250		H1 73 320
33 411429.8 dec 02505704	66 230	22 2/46334	H1 72 204
	63 553	22 4344936	H1 75 257
	64 553	32 411429.8.dec 3369795	H1 75 201
22 123.0.080 92354084 1	69 553	30 411429.8.dec 6106562	H1 76 204
22 44 45.0.000 9359/881 1	70 553	711449.8.dec 2742002	114
22 7723.0.0ec 983855H1 1	71 439	411429.8.dec 6160891	
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33 411429.8.dec d2350550		33 411429.8.dec 4047691	
33 411429.8.dec 03022570			H1 72 360
33 411429.8 dec 03804700		33 411429.8.dec 660714H	72 340
33 411429 8 dec =2700000		33 411429.8.dec 15447631 33 411429.8.dec 15447631	11 72 156
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20 1117E3.0.000 4/50950H1 20	6 447	411429.8.dec 142400cT	
33 411429.8.dec 3818447H1 20		711429.8.Qec 4566527L	
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411429.8.dec 21447144 12			,
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33 411429.8.dec gazagaa			72 335
33 411429 8 dec 200010011	564	32 4425.0.dec 3900464H	1 72 220
33 411429 8 dog 500 4150 415	293	32 411429.8.dec 4045408H	1 100
32 4445H1 36	292	30 411429.8.dec 1730041H1	1 004
32 411429.8.dec 583800H1 38	000	11429.8.dec 173004+E6	
33 411429.8.dec g1390456 34		33 411429.8.dec 3440050U4	
411429.8.dec 3143351H1 05	^==	33 411429.8.dec 187052414	
411449.8.dec 6360697U4	377		
33 411429.8.dec 02307550	374		
33 411429.8.dec 498375744 210	554	5945503H1	72 365
33 411429 8 dec 2000000111	40-	2 11425.0.0EC 4001455H1	72 346
20 1-0.0.dec 3900386H1 241	E10	2728288H1	72 316
22 THE STATE OF TH	504	3832210H1	
22 411429.0.dec 4500345H1 240	540	411429.8.dec 132200744	
33 411429.8.dec 4500394H1 249		411429.8.d c 227042214	72 304
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34	320674.7.d	C 03050504		i abie	2 cont.		,	1 01/03	00/25610
34	320674.7.d		575	678	34	200074			
34	320674.7.de		583	671	34	320674.7.dec	2783335	H2 34	070
34	320674.7.QB(c 1382266H1	333	577		320674.7.dec	7005400		278
34	320674.7.ded		333	596	34	320674.7.dec	72177611		258
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34	320674.7.d c	01766401		622	34	320674.7.dec		H1 34	258
34	320674.7.dec	g2563628	351	588	34	320674.7.dec		1 34	282
34	320674.7.dec		184	518	34	320674.7.086		11 34	360
34	320674.7.dec		251	614	34	320674.7.dec	g1775483	37	155
34	320674.7.080		254	615		320674.7.dec	5033665H	11 34	
34	320674.7.dec				34	320674.7.dec	4844745H		295
	320674.7.dec	4336444		620	34	320674.7.dec	2659859H	47	272
34	320674.7.dec	03753004		620	34	320674.7.dec	1000040		288
34	320674.7.dec	g1933057		622	34	320674.7.dec	1922312H	1 . 47	316
34	320674.7.dec		242	615	34	320674.7.0BC	6169157H	1 46	136
34	320674.7.dec	4914556T8		573	34	320674.7.dec	6008947H	1 62	371
34	320674.7.080	2324436H1		510		320674.7.dec	6508822H	1 65	
34	320674.7.dec	2323128H1		506	34	320674.7.dec	2174565H1		157
34	320674.7.dec	g1898237			34	320674.7.dec	2922846H1		198
	320674.7.dec	1910935H1		515	34	320674.7.dec	3720640114	68	336
34	320674.7.dec	92195148		501	34		3739613H1	68	247
34	320674.7.dec	4358201H1		520	34		3175571H1	68	321
34		3304707H1	48 3	30	34		1623101H1	46	256
34		3391767H1	49 3	30	34	320074.7.Gec	5289080H1	46	205
		g1795515		03		3400/4./.dec	638816H1	46	
34	320674.7.dec	5345961H1		90	34	320674.7.dec	1894775H1		190
	3206/4./.dec	1748443H1			34		1748326H1	46	296
	9200/4./.dec	E160000	• • • • • • • • • • • • • • • • • • • •	84	34		5554007U	52	289
34 (320674.7.dec	2225		52	34		5554087H1	52	318
34 3		71000474	776 11	153	34		167107H2	54	275
34 3		~7^^~	789 10	83			46643H1	54	280
34 3		766921	793 10	16	34	320674.7.dec 1	705647H1	54	
34 3		2479851H1		88	24	3400/4./.dec 3	560375H1	56	286
34 3	20674.7.dec 4	746673H1	26 25		34 (24V0/4./.dec 3	834716H1		335
34 3	200/4./.dec 4	848226H1		^	34 3	320674.7.dec 1	972338H1	57	237
	<00/4./.dec 4	4000000			34 3		E12000H1	57	299
34 3	20674.7.dec 2	^^======			34 3		513638H1	57	294
34 32	20674.7.dec 2	007466	5 41	7 ;	34 3	20674.7.dec 58	857125H1	58	331
34 32		140700		3	_		356825H1	58	317
34 32		118730H1 2	9 313	•	_	20074.7.dec 60)27442H1	58	353
34 32		2001453 7	3 380		- J	400/4./.dec 63	21410H1	59	
34 32		570374H1 7	4 191		34 3	<ud>028</ud>	15776H1	_ = =	346
	00/4./.dec 14	142776H1 7			94 3	20674.7.dec 50	47756H1		328
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34 32	V0/4./.dec 12	ACCOALL			4 32		2090/H1	60	299
34 32	0674.7.dec 10	400000		3	4 32	20674.7.dec g2	92839H1	592	832
. 34 32	0674.7.dec 10.	42000114		3	4 32		616212		1129
34 320		COCCCI		3.	4 22		57122H1		906
34 320		69622H1 14	3 381	3.	7 OZ	06/4./.dec 64	1657H1	`	955
34 320		260477 14				VD/4./.dec ass	53248		
		753672 15		34	32	0674.7.dec 220	5827H1		005
04	0674.7.dec 383	39247H1 150		34	32	0674.7.dec 334	2350H1	'	019
	/º/4./.dec asc	60404 16		34	320	0674.7.dec 371	7588H1		93
	10/4.7.dec 050	38447 182		34	320		535H1	32 3	26
34 320	0/4.7.dec naa	144004		34	320		^	32 1	41
34 320	674.7.dec 275	740011		34	320		67659		07
34 320		707111		34	350		5163H1		71
34 3206		700		34	220	674.7.dec 511	8730F6	29 4	
34 3206		73968 264	555		320	0/4./.dec 497	3794H1	'`	
34 3206	574.7.dec g418	86990 265		34	320	674.7.dec 5564		-·	
	74.7.dec g290	01581 266		34	320	674.7.dec 5403	1050114	- <i>-</i>	
	7/4.7.dec 20ar	2000111	615	34	320		FACA	31 26	9
<i>3</i> 4 3206	74.7.dec 6168	2400111	539	34	320		5030	32 3 ₆	1
34 3206		2007	615	34	3304		545H1 3	31 28	
34 3206	74.7.dec 0225	4000	615	34	3200	574.7.dec 4975	303H1 3	7 31:	
34 3206			615	34	3200	2/4./.dec 1118	46R1 3		
34 3206		954H1 39	270		3206	74.7.dec 1118	46R6 3	_ •	
	74.7.dec 2470	102H1 30		34	3206	74.7.dec 3100	700114		
	/4./.dec g128	1396 42	245	34	3206	74.7.dec 0103			
34 32067	(4.7.dec 285nd		436	34	3206				
<i>3</i> 4 32067	74.7.dec 4055	446114	352	34	3206		63H1 3	9 299	
34 32067	4.7.dec 58672	200114	162	34	3200		323H1 1		
34 32067		303H1 39	321	34	3200	74.7.d c 26359	90H1 2	250	
34 32067		33H1 45	336	34	SZU6	(4./.dec 53498	41H1 12		
		154H2 A1	327		3206	/4./.Dec 1110 <i>a</i>			
	4.7.dec 42752	74H1 45		34	32067	74.7.dec 20077	4		
√- 320574	4.7.dec 25224	62H1 34	328	35	19726	7.1.dec 14720	00114	287	
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				88		7.1.dec g1548	716 7	330	
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	-4040204	1	464	35	19726		4567732H1		759	
35 ` 1	197267.1.dec 915-0204	40	222	35	19/20	57.1.dec 57.1.dec	374248H1	498	720	
35		40	208	35	19720	67.1.dec	1473995H1	616	805	
		40	370	35	1972	67.1.dec	5986850H1	620	901	
_	197267.1.dec g1859154 197267.1.dec 4772879H1	348	629	35 35	1972	67.1.dec	a4511097	621	901 901	
	107267 1 dec 319110T6	345	865	35	1972	67.1.dec	6212219H	1 625	914	
	107267 1 dec 5028542H1	423	679	35	1972	67.1.dec	g4332172	627 6 636	862	
35 35	107267 1 dec @2537655	437	871 574	35	1972	67.1.dec	2132792T6	•	527	
35 35	407067 1 dec 9///9101	357	605	35	1972	67.1.dec	3374148H g1025985		535	
35	197267.1.dec 424019011	359 361	606	35	1972	267.1.dec	g896704	276	572	
35	197267.1.dec 2//1133H	357	785	35	1977	267.1.dec	2751121H	11 551	642	
35		373	512	35		267.1.dec 267.1.dec	319110H1	228	610	
35		461	913	35		267.1.dec	46510301	11 220	462	
35	197267.1.dec g3254447 197267.1.dec g3961279	466	901	35 35		267.1.dec	3246169h	41 221	454 492	
35	107267 1 dec 1637850H1	450	654	35	197	267.1.dec	48505021	41 213 6 228	742	
35 35	107267 1 dec 1348899H1	438	697 654	35	197	267.1.dec	3191100	_		
35 35	197267.1.dec 1635340H	450	531	35	197	7267.1.dec	13489/0			
35	197267.1.dec 914/115/	339 403	565	3	5 197	7267.1.dec			931	
35	197267.1.dec 241314501	427	521	3		7267.1.ded		T6 678		
35	197267.1.dec 612807H1 197267.1.dec 5122537T6	413	870	3		7267.1.de 7267.1.de		684		
35		418	644			7267.1.de	c 2103287	7H1 690	- 700	
35	197267.1.dec 5213230H1 197267.1.dec 4365757H1	421	681		5 19 5 19	7267.1.de	c 2872292	2H1 46		
35	107267 1 dec g896705	691	905		15 19	7267.1.de	C 4950/30	0H1 469 3H1 45		
35 35	107267 1 dec 03601113	695	- 40		15	7267.1.de	C 326/6/			
35	197267.1.dec 2/51946F1	389 386			35 19	7267.1.de	ec g511023 ec 537271	,		4
35	197267.1.dec 192760171	70€		;	35 19	97267.1.de			1 85	
35	197267.1.dec 5551489H1 197267.1.dec g1855688	706				97267.1.de 97267.1.de	ec 398676	19H1 20		
35	G2464352	75				97267.1.d	ec a28360)64 /	79 90	
35	107067 1 dec . 01471158	75		•	35 1 35 1	97267.1.d	ec a36657	793 /	90 00 90	
35	407267 1 dec 214/0/01	76			35 1	97267.1.d	ec gasoar	091 8	00 90 26 61	
35 35	107267 1 dec g1114641	69			35 1	97267.1.0	lec 25003	70. U		54
35	197267.1.dec 611349F1	56 1 56			35 1	97267.1.0	iec 388642 iec 26045		13 3	70:
35	197267.1.dec 2491963H	, 56 56				197267.1.0		01H1 2		01:
35	197267.1.dec 93232143	57				197267.1.0 197267.1.0	dec 27171	83H1 1		59
35	1548831	58	34 87		35 35	197267.1.	dec 36146	68H1 1		17 80.
3!	5 107267 1 dec 5164893H	1 59	99 86		35	197267.1.	dec 2500 l	10 11 7 1		101
3	5 107267 1 dec 817768H1	3	17 56 76 51	_	35	197267.1.	dec g3//	<i></i>		361
	197267.1.dec 2501176F		76 51 78 52		35	197267.1.			977	1229
	197267.1.dec 9842912		23 56		36	332335.1 332335.1		50H1		1007
3			31 85	. 8	36	332335.1	dec 4556	145H1		1173
	1803588	r6 3		63	36 36	332335.1	dec 3430	852F6	•	1428 1268
	107267 1 dec g116613	3	395 56		36	332335.1	.dec 1335	220H1	1034 1949	2100
	107267 1 dec 0536501	3 :		13 01	36	332335.1	.dec 418/	132H1	1993	2225
	35 197267.1.dec 9400535	5 3		03	36	332335.1		0021H1 3581H1	2062	2315
	35 197267.1.dec 5992654			02	36	332335.1		57941	2073	2387
			378 5	96	36	332335. 332335.	1 dec 487	5910H1	2074	2367
			227 7	115	36	332335.	1 dec 341	4856H1	2180	2430
	35 107267 1 dec 5313986	iH1		179	36 36	332335.	1 dec 489	6022H1	1994	2275 1756
	35 197267 1 dec 3984239	9H1		539 601	36	332335.	1.dec 645	7742H1	1201 1363	1550
	35 197267.1.dec 9810074	2		723	36	332335	1.dec 9/2	3998 3966F6	1382	1883
	35 197267.1.dec 920044	05 07		919	36	332335		34333F6	614	1169
	35 197267.1.dec 939860	3/ LI4		796	36	332335		64333H1	614	883
	35 197267.1.dec 906113	R1	541	901	36	332335 332335	1 dec 32	12701H1	1	293
	33 13770	2H1	436	685	36	332335	1 dec 64	21135H1	202	783
	35 055324	H1	438	680	36 36	33233	1 dec 33	78675H1	388	636 1724
	of 107267 1 d c 955324	IR1	438	901	36 36	33233	5.1.dec g7	51133	1469 1567	1771
	35 107267 1 d c 180358	38H1	72	335 277	36	33233	5.1.dec 49	33390H1	1676	
	35 197267.1.d c 14940	78H1	66 119	392	36	33233	5.1.dec 18	305556F6 305556H1	1676	1931
	35 197267.1.dec 22//9	53H1 40⊔4	126	376	36			773837H1	1716	
	35 197267.1.dec 25003	30H1	453	722	36	33233	5.1.dec 37	, , 050		
	35 197267.1.dec 48353	JU: 11			89					

PCT/US00/25610 Table 2 cont.

36 332335 1 dog 47000		Table 2	cont.	i	PCT	/US00/25610	
36 002000.1.060 4/660	37H1 1849	2127					
36 20000.1.080 53945	18H1 1375	1544	37	238992.13.dec 57842	262H1	760 00.	
36 302000.1.0 € 15809	56H1 1382	1579	37	~~~~~~~. IS.DBC 03425	2016	760 991 763 1145	
36 50000.1.dec 611517	74H1 4na	652	37	200332.13.0ec a1854	CCC	40)
36 332335.1.dec 472417	78H1 498	754	37	200332. IS.DEC 90555	OLI4		
36 332335.1.dec 309615	9H1 1460	1730	37	200332.13.08c 16352	ATU	645 937	
36 332335.1.dec 369142	9H1 530	749	37	~00332. IS.Dec narro	620	734 938	
36 332335.1.dec g72391	3 2634	2724	37	200332.13.dec 10863	ACTA	737 1147	
36 332335.1.dec 234119	H1 2515	2722	37	-00334. IJ.dec 57925/	ROU4	764 1166	
36 332335.1.dec 184386	6H1 2588	2732	37	200332.13.0ec 17017/	אדבר	760 1057	
36 332335.1.dec g22386	87 2504	2698	37	230332.13.dec 232343	20L4 .	639 1164	
332335.1.dec 175141	ELI4	2724	37	200332.13.dec 782457	' ⊔∢ .	643 897	
30 332335.1.dec 175141	EC	2698	37	238992.13.dec 228032	T1 (636 882	
35 332335.1.dec 1751416	70	2698	37	238992.13.dec 277689	8H1 (636 902	
36 332335.1.dec 343095	TC	2698	37	238992.13.dec 212926		537 900	
30 332335.1.dec 1805556	TC CO	2725	37	238992.13.dec 560897	9H1 1	053 1132	
30 332335.1.dec 0751124		2730	37	238992 13 dec 540897	H1 1	075 1138	
37 238992.13.dec 4801804	LI4 - 1	2724	37	238992.13.dec 543395	4H1 1	091 1207	
- EUUSSE, 13 DEC 0/72000	•	270	37	238992.13.dec 3318824	1 H1 7	63 1032	
- 430332.13 dec 53350cc	7 844	1138	37	238992.13.dec g375317	70 7	65 1147	
37 238992.13.dec 1981188		555	37	238992.13.dec g227745	:4	70 1062	
37 238992 13 doc 20000	H1 385 5	566		230332.13.0ec 5420344	LI4 A	11 823	
37 238992.13.dec g2000913	416 5			200554.13.0ec 0110307	·c -	10 799	
37 238992.13.dec 506041H				EGGGGG, IG. GEC GEAGAAA	LI4 A.		
THE PROPERTY OF THE PROPERTY O			••	200992.13.dec 3385322	LI4		
		~~		200394.13.dec 2800701	U1 ~	_	
	14 40-			-50332.13.dec 29511011	U4		
200332.13.08C 4132042L	14 00-	~ ~ '	- 4	-00332.13.08C 1300020	TC 00		
200332.13.00C 2226878T	6 000	• • • • • • • • • • • • • • • • • • • •		-90334. IJ.DEC 3843600L	4		
200332.13.0ec 4131050L	2 222			-00334.13.dec n375606:			
230392.13.dec 5911619L	1 010	` `		-00002.13.0ec 6107000L	14		
200332. J3.Gec 2262124D	,	`	- ح	238992.13.dec 4847550H	11 65		
230392.13.0ec 994076114			37 2	38992.13.dec 5330638T	-		
200332, I3.0ec 18110core			7 2	38992.13.dec g2752175	6 668	3 1109	
200992.13.dec 7810/2014		00 з	7 2	38992.13.dec 1552390H	671	838	
230392.13.dec 2182095U4	636 91		7 2	38992 13 dec 2001	1 680		
200332. IS.GEC 610504U4			7 2	38992.13.dec 2081835T	683		
230332. 13 dec 520400014	170 42	7 3		38992.13.dec 5606465H	1 682	910	
37 238992.13.dec 921438H1		7 37		38992.13.dec 1473638H	1 645		
37 238992.13.dec 6495685H1	222 499	37		38992.13.dec 1655911H			
37 238992.13.dec 2465087H1		37		プララム・13.0ec 1798756Td	506	1098	
37 238992.13.dec 5870435H1	237 430	37	20	70332. IJ.Dec 506181114	400	672	
37 238992.13.dec 4466185H1	239 527			10332. 13.0ec 1214020114		706	
37 238992 13 dos =24466185H1	263 436	37	23	^{©33} 2.13.08C 4072200⊔∢		1098	
37 238992.13.dec g3148442 37 238992.13.dec g3148442	278 680		23	0332.13.0ec 5680227L14		1104	
-50002.13.UEC 180007004	335 583	٠,	20	0992.13.dec 418521701	870	1104	
	602 886	0,	200	⁰³³ 2. I3.0ec 817636⊔4	881	1205	
	601 853	37	230	9992.13.dec 6102144U4	883	1138	
	601 811	37	200	2332.13.0ec 817626T4	882	1137	
	831 1132	37	200	2226. IJ.DBC 3320104TA	887	1104	
200332.13.0AC 66/207114	000		200	7554.13.08C 3621515U4		1169	
230332.13.0ec 03755470	838 1109 840 1139	•	200	1334. 13.0ec a825072	888	1105	
200332, 13.dec 5222770U4	244	•	430	1992.13.dec 0678520	890	1148	
200992. IS.Dec 314381414	844 1018 823 1141	•	230	332.13.0ec 3458087U+	890	1138	
200392.13.0ec 31/3107U4	000		230	334. IJ.Dec 05150240	891	1138	
200332. I3.0ec 634398EU4	CO-	37	200	334. IJ.Dec 3400541114	913	1138	
200332.13.08C 65381/6U4	504	37	200	794.13.0ec 2560561U1	915	1138	
200332.13.0ec 1086306U4	500	37	2389	992.13.dec 4368210H1	919	1178	
200992.13.0ec 5313607U4	539 801 539 789	37	2389	992.13.dec 2357876H1	942	1138	
200332. IS.Dec 327020EU4	COO	37	2389	92 13 dos 2700 5	945	1138	
200332.13.0ec 198620cpc	539 765	37	2389	992.13.dec 3729584H1	947	1138	
200332.13.0ec 767764Te	539 910	37	2380	92.13.dec 4509315H1	955	1231	
230332.13.dec 1637074114	542 1093	37	2380	92.13.dec 4542916H1	955	1225	
- 200332. IS DEC ARRODOMINA	577 806	37	2300	92.13.dec g706241	965	1138	
37 238992.13.dec 1655010T6	582 880	37	2300	92.13.dec 788265H1	986	1138	
37 238992.13.dec g907966	580 1174	37	2305	92.13.dec a2215/as	1025	1148	
37 238992 13 doc 441050000	595 794	37	2303	92.13.dec 3150/15L14	822		
	718 1104 .		2303	24.13.0ec 1340465U+	~~ .	1115	
	698 749	37 37	20033	74. IJ.DEC 51028664	~	981	
200332. 13.0ec 05546066	728 1139	37 37	£3033	74. IJ.Dec 3860027⊔4		1037	
- £30332.1.1 dec 11ecco	711 865	37	E0033	/く・/ J.Dec 632725204		1061	
	. 003	37	23899	2.13.dec g3092108		1042	
		90		3100	795	1137	



				Tab	le 2 cont.	•		1,0300/	
38	199736.1.dec	g2903286	274	597	41	481454.4.dec	719870H1	1007	1256
. 38	199736.1.dec	1522915H1	380	484	41	481454.4.dec	965365R1	1028	1602
38	199736.1.dec	6593543H1	423	940	41	481454.4.dec	965365H1	1028	1348
38	199736.1.dec	2890957H1	669	931	41	481454.4.dec	1422810H1	1031	1281
38 38	199736.1.dec 199736.1.dec	g2432628 g1967342	1	333	41	481454.4.dec	705397H1	166	315
38	199736.1.dec	g1967342 g3751279	1 1	450 444	41 41	481454.4.dec		179	342
38	199736.1.dec	1444281H1	1	257	41	481454.4.dec 481454.4.dec	712266H1 3442620H1	181	277
38	199736.1.dec	1444281F6	i	505	41	481454.4.dec	724216H1	186 1720	441 1958
38	199736.1.dec	g4289858	4	321	41	481454.4.dec	724216R6	1720	2130
38	199736.1.dec	1439532H1	13	297	41 .	481454.4.dec	551787H1	1737	1894
38	199736.1.dec	1439532F1	13	501	41	481454.4.dec	722578H1	1738	2006
38	199736.1.dec	g2457237	18	476	41	481454.4.dec	1423789H1	1527	1740
38	199736.1.dec	g3043040	18	436	41	481454.4.dec	1385707H1	1538	1767
38 38	199736.1.dec	3034279H1	58	354	41	481454.4.dec	1388173H1	1538	1785
39	199736.1.dec 228864.5.dec	3031157H1 943941T1	58 417	360 714	41	481454.4.dec	4356537H1	1542	1660
39	228864.5.dec	2240562H1	433	692	41 41	481454.4.dec 481454.4.dec	1261250H1 1261250R6	1542	2131
39	228864.5.dec	2569601H1	437	681	41	481454.4.dec	1261250H1	1542 1542	2051 1774
39	228864.5.dec	751050H1	466	691	41	481454.4.dec	4755269H1	1597	1863
39	228864.5.dec	4424104H1	493	743	41	481454.4.dec	3553105H1	1603	1900
39	228864.5.dec	2101758H1	516	757	. 41	481454.4.dec	3741144H1	1646	1862
39	228864.5.dec	5608727H1	526	747	41	481454.4.dec	698444H1	1650	1893
39	228864.5.dec	137261H1	217	384	41	481454.4.dec	1423073H1	1667	1876
39 39	228864.5.dec 228864.5.dec	3469340H1 2438390H1	217	460 456	41	481454.4.dec	2497753H1	1678	1923
39	228864.5.dec	2438390H1 2438735H1	224 224	456 437	41 41	481454.4.dec	722589H1 725929H1	1801	2056
39	228864.5.dec	4839445H1	263	537	41	481454.4.dec 481454.4.dec	725929H1 710840H1	1808 1834	2056 2089
39	228864.5.dec	4698944H1	319	576	41	481454.4.dec	6554952H1	1852	2375
39	228864.5.dec	503758H1	322	558	41	481454.4.dec	192279F1	1854	2273
39	228864.5.dec	516461H1	383	598	41	481454.4.dec	1262802H1	1370	1565
39	228864.5.dec	2413301H1	387	600	41	481454.4.dec	722414H1	1357	1609
39	228864.5.dec	873275T1	414	714	41	481454.4.dec	1262803H1	1371	1564
39 39	228864.5.dec 228864.5.dec	873275H1	414	652	41	481454.4.dec	727709H1	1400	1616
39	228864.5.dec	g2166848 6513978H1	1	514 543	41 41	481454.4.dec	368161H1	1410	1671
39	228864.5.dec	3321280H1	i	259	41	481454.4.dec 481454.4.dec	3248182H1 192279H1	1469 1490	1738 1712
39	228864.5.dec	3537292H1	4	270	- 41	481454.4.dec	192279R1	1490	2004
39	228864.5.dec	3974316H1	4	291	41	481454.4.dec	719958H1	1502	1720
39	228864.5.dec	6269744H1	15	506	41	481454.4.dec	710004H1	1504	1741
39	228864.5.dec	2276784H1	27	263	41	481454.4.dec	1386020H1	1525	1696
39	228864.5.dec	2707203H1	175	414	41	481454.4.dec	720247H1	1527	1699
39 39	228864.5.dec 228864.5.dec	4317110H1 5106358H1	176	452	41	481454.4.dec	1423446H1	320	467
39	228864.5.dec	1886806H1	184 214	430 487	41 41	481454.4.dec	3214020H1	320	574
39	228864.5.dec	g1576711	215	374	41	481454.4.dec 481454.4.dec	2991370F6 1387254H1	318 320	524
39	228864.5.dec	604541H1	217	436	41	481454.4.dec	1387640H1	317	529 456
40	986539.1.dec	g4085132	256	725	41	481454.4.dec	2991370H1	318	557
40	986539.1.dec	94222648	263	598	41	481454.4.dec	712619H1	191	315
40	986539.1.dec	g865178	246	530	41	481454.4.dec	3346330H1	312	593
40	986539.1.dec	6432233H1	247	648	41	481454.4.dec	1261236H1	317	566
40	986539.1.dec	g1064176	290	601	41	481454.4.dec	1261236R1	317	795
40 40	986539.1.dec 986539.1.dec	g3053295	238	600	41	481454.4.dec	6316533H1	1	301
40	986539.1.dec	g2912341 g2902568	305 310	686 466	41 41	481454.4.dec 481454.4.dec	4773101H1	2287	2413
40	986539.1.dec	g1757251	1	415	41	481454.4.dec	3388764H1 3738442H1	1107 1747	1386
40	986539.1.dec	g4888422	1	226	41		1363446F1	1763	1982 2199
40	986539.1.dec	g2898861	314	596	41	481454.4.dec	1363446H1	1763	2019
40	986539.1.dec	g1678784	547	937	41	481454.4.dec		1766	2003
40	986539.1.dec	6160421H1	549	835	41	481454.4.dec		1770	1882
40	986539.1.dec	g2526384	20	245	41	481454.4.dec	1422809H1	1031	1280
40	986539.1.dec	493287H1	202	416	41	481454.4.d c	3550570H1	1045	1177
40 40	986539.1.dec 986539.1.dec	4433535H1	640	912	41	481454.4.dec	3553072H1	1063	1360
40	986539.1.dec	g2714767 g2703735	732 789	1240 1257	41	481454.4.dec		1106	1229
41	481454.4.dec	3552820H1	1124	1257 1408	41 41	481454.4.dec 481454.4.dec	2495808H1 708530H1	1863	2212
41	481454.4.d c	4239624H1	980	1237	41	481454.4.dec	700530H1 720746H1	1874 1869	2093 2087
41	481454.4.dec	727425H1	998	1225	41	481454.4.dec	719117H1	1919	2136

	0 01/2	23330	1									
					Tal	ole 2 co				PCT/U	JS00/256	10
	41 48145	4.4.dec 724	1019H1	1933								••
	41 48145	4.4.dec 138	8274H1	1942	2141		42 47480	0.7.dec	336413	2L14 .		
	41 48145 41 48145	4.4.dec 719)545H1	1949	2188 2145		42 47480	0.7.dec	6206404			06
	10170	4.4.dec g83	1279	2034	2426		42 47480	0.7.dec	0750600			93
	.0,175	4.4.dec g89	1552	2051	2395		42 47480	0.7.d c	0226074	- ·		81
	.0.170	4.4.dec g64	6749	2091	2375		47480	0.7.dec	4800060			42
	.0170-	4.4.dec g20	17804	2144	2380		2 47480	0.7.dec	5866864	H1 2		
	101707		5110H1	2201	2372		2 474800 2 474800	0.7.dec	4974548	H1 2	85 45 87 55	
	41 481454 41 481454		2647H1	750	980			7.7.dec	671748H	1 . 2	88 53	
	41 481454		2647R6	750	1196	4.).7.dec	4626725	H1 20	90 55	
	41 481454		542H1	899	1196	4:		7./.dec	45499371	H1 20		
	41 481454		30R6	961	1399	42		./.dec	702403H	1 20		
	41 481454.		30H1 92F1	961	1242	42	474800	.7.dec	3085429	11 29	2 574	
	⁴¹ 481454.	4.dec 1902	92F	976	1504	42	474800	7 dec	2536581F		7 554	
	⁴¹ 481454.	4.dec 1902	32MI 92D1	371	594	42			4577417H			3
	⁴¹ 481454.	4.dec 1389	249H1	372	916	42			3278477H			,
	⁴¹ 481454,	4.dec 2987	096H1	415	641	42	474800.	7.dec	3525826H 2274935H		_ +00	
	41 481454,4	4.dec 0573/	061		526	42	474800.	7.dec	6128496H			
	481454.4	4.dec 72163	32H1	4	781 704	42	474800.1	7.dec	3538467H			
	11 481454.4 11 481454.4	4.dec 58966	60H1		734 816	42	474800.7	7.dec (g1696705			
		1.dec 40284	67H1	`	B31	42	474800.7	7.dec	4654475H	255 1 254		
			08H1	`	915	42	474800.7	dec 2	2519031H	1 253	,	
	1 481454.4 1 481454.4		36 (922	42 42	474800.7	dec 1	1415905H1	253		
4	.0,404,4		07H1		009	42	474800.7	odec g	11991337	254		
4				⁷ 50 1	243	42	474800.7	.dec 6	3128132H1	256		
4	481454.4			163 1	277	42	474800.7 474800.7	.dec 6	i512254H1	256	764	
41	481454.4.	.dec 664692 .dec 373953			371	42	474800.7		61646T1	255	739	
42	474800.7.	dec nanese		345 1	520	42	474800.7		127445H1	256	680	
42	474800.7.	dec 433301			90	42	474800.7.		921246H1	254	546	
42	474800.7.	dec dannes			95	42	474800.7.		754734H1	255	486	
42	474800.7.6	dec 031547			96	42	474800.7	dec 31	154991H1 162126H1	257	559	
42	474800.7.	dec 137656	7F1 3			42	474800.7.	dec 32	135104H1	255	531	
42 42		dec 386604	6H1 3/	50 78 50 63		42	474800.7.6	dec 51	70106H1	255	529	
42		dec 137656	7H1 36			42	474800.7.0	dec 51	18447H1	256 256	534	
42		ec 445465	7H1 36	57 56		42	474800.7.c	dec 34	93016H1	255	521	
42	474800.7.d 474800.7.d		36 36			42	474800.7.0	iec 35	51427H1	255	348 546	
42	474800.7.d		'4 37	0 78		42 42	474800.7.d	rec 31	48115H1	255	542	
42	474800.7.d		1 37			42	474800.7.d	lec 33	18467H1	252	513	
42	474800.7.de				3	42	474800.7.d	ec 22	71268H1	256	517	
42	474800.7.da	BC 3436007	114 =			42	474800.7.d 474800.7.d		48927H1	255	477	
42	474800.7.de	C 20470co				42	474800.7.de		1653H1	255	491	
42	474800.7.de	C 3471500				42	474800.7.de		16336H1	256	525	
42	474800.7.de	C 2005746				42	474800.7.de		35584H1	257	518	
42	474800.7.de	C 0205200	11 261 7 262			42	474800.7.de		6988H1 0527H1	257	458	
42	474800.7.de	C 43006201	11 264			42	474800.7.de	C 037	36691	255	554	
42 42	474800.7.de	C 2452854H	11 263			42	474800.7.de	K 490	3583H2	736	786	
42	474800.7.de	c 2365057h	11 263			42	474800.7.de	c 337	5214H1	619 631	771 700	
42	474800.7.dec		11 263			42	474800.7.de	c g27;	37517	634	789 780	
42	474800.7.dec		1 263	444		42 . 42 .	474800.7.de	c g21(84209	657	789 789	
42	474800.7.dec			509			474800.7.de	c g218	84197	658	789	
42	474800.7.dec			505			474800.7.dec		38940		790	
42	474800.7.dec	3968293H 575286H1		-538			174800.7.dec		668H1		787	
42	474800.7.dec		264	564			74800.7.ded 74800.7.ded		6489		789	
42	474800.7.dec	01642050		499			74800.7.dec			684	796	
42	474800.7.dec	31820161	266	548		•	74800.7.dec			683	789	
42	474800.7.dec	497974611		575		42 4	74800.7.dec		~~~	687	789	
42	474800.7.dec	g1698167		534			74800.7.dec			699 7	789	
42	474800.7.dec	1467071H1	259 250	567	4	12 4	74800.7.dec				785	
42	474800.7.dec	g1067573		457	4	12 4	74800.7.dec	g2261 g4075			789	
42	474800.7.dec	4556026H1	261 261	490		2 47	74800.7.dec	g1148			'88	
42	474800.7 dec	2367365H1	261 260	520		2 47	74800.7.d c	40126		732 8	10	
42	474800.7.dec	1004773H1	260 261	497		2 47	4800.7.dec	21179	100114		32	
42 4	474800.7.dec	1493863H1	283	448 506		2 47	4800.7.dec	56344			54	
76	474800.7.dec	g894785	284	381		2 47	'4800.7.dec	25340	90H1 ^		78	
			-07	20 [4:	2 47	4800.7.dec	27148			B0	
					92				1 2	JO 46	65	

		TC1.00	
WO 01/23558	Table 2	nont	
110	lable 2	2076719H1 2/0	
	242 481	42 474800.7.dec 3370713.1.	
42 474800.7.dec 2210110H1		42 474800.7.dec 90-77.	
474900 7 dec 221599501		42 474800.7.dec 94629546 551 789	
474900 7 d c 730002R1	244 587	40 474800 7 dec 0113/292	
42 Jan 7 doc 3566922H1	239 451	45 474800 7 d c 6296919H1 400	
	244 549	42 3 400 2397267 480 773	
474800.7.dec 371400077	244 488	42 7 dee 9389445 489 775	
474800.7.dec 91954050	247 706		
474900 7 dec 7499/501		42 474800.7.dec 375000 503 788	
1259/20F1		42 474800.7.dec 6348/2011 515 797	
42 4834575H1	248 515	42 474800.7.dec 12356/6F1 515 791	
42 730002H1	244 464	474900 7 dec 12350/001	
	244 460	474000 7 dec 52680/UH	
	246 501	42 7/100 2884366H1 555 /89	
42 474800.7.dec 451427711	246 721	42 1372524H1 556 //9	
42 474800.7.dec 2494966F6	246 506		
474800 7 dec 2149200011	405		
474900 7 dec 2729518011	400	42 474800.7.dec 4115160111 525 773	
474900 7 dec 2211220H1	246 488	474900 7 dec 13498/4FT	
42 1696726H1	235 449	474900 7 dec 526976001	
	239 433	474900 7 dec 2752//9H1	
	246 475	42 474000 7 dec 0895371 558 700	
	545	72 7 dec 94147221 545	
42 474800.7.dec 495752011	500		1
474800 7 dec 2893665FT	540		5
474800 7 dec 22088940	500	42 474800.7.dec 155005.11 540 78	7
474900 7 dec 480330/T	248 508	42 474800.7.dec g1/28042 603 78	
42 Jan 2480259H	1 240 4/2	474900 7 dec 08129/4	
42 402691H1	246 320	474900 7 dec 249490010 000	
	1 247 508	474900 7 dec 02 104440	
	1 247 527	42 7 7 7 7 7 7 7 184427 611	
		72 -1000 7 doc g4149309 61/	39
474800.7.dec 13573201	500		30
42 474800.7.dec 4952010F			B2
474800 7 dec 1918861F	540	42 474800.7.dec /49370114 047 4	74
474900 7 dec 662273H	248 512	42 474800.7.dec 1260232111 247 4	60
42 7 400 1357326	11 248 400	474900 7 dec 1259/2501 24	43
42 5669715	11 244 42!	474000 7 dec 2906023F1 240	14
	1 24/ 455	474800 7 dec 3136046H1 248	
	H1 232 422	42 3401103H1 248	198
42 474800.7.dec 1591050			194
42 474800.7.dec 1288010	400	42 49 4920352H1 249	520
474900 7 dec 1286624		42 474800.7.dec 4020001114 047	500
474900 7 dec 4731307			495
474900 7 dec 3/49043	IN	42 474800.7.dec 3297623111 348	499
474900 7 dec 1922474	H1 280 517	474800 7 dec 1211350H1	490
42 THORS 7 des 1340372	9H1 283 324	474900 7 dec 52/922001	533
	9H1 283 300	474900 7 dec 371922501 245	491
	5H1 235 456	42	
4/4000		42 4068071H1 248	509
42 474800.7.dec 129422	001		495 .
474800.7.dec 913241	45 25	4/ 7/7000	364
42 474800.7.dec 129513		47 474000.7.000 0= 040	498
474900 7 dec 380550	9111	42 474800.7.dec 4963531111 248	473
474900 7 dec 46630	OLIT TO	42 474800.7.dec 2/23303111 076	540
474800 7 dec 237287	9H1 23/ 40	42 474800.7.dec 61/5/31H1 276	526
42 445984	10H1 232 40	474900 7 dec 92441201 270	567
42 4740007 400 40206	79H1 234 ³⁰	* = 474000 7 dec 42/403901 2/0	
	55H1 235 48	2 42 This 7 day 4158452H1 2/6	507
		2 42 7 400 9834460 2/6	549
42 474800.7.dec 31400	DUN1		505
474800.7.dec 32/01	40111 202		382
474800 7 dec 2501/	03111	474800.7.dec 392545711	596
474900 7 dec 58497	17H1 234 5	42 474800.7.dec g203/450 277	790
42 THOSE 7 dec 12944	27H1 234 4	474900 7 dec 22/48201	651
42 Trees 7 dec 26072	69H1 235 4	9 42 474800.7 dec 924412R1 2/6	
	02H1 232	39 42 474900 7 dec 989831H1 2/6	497
		36 42 474000 7 dec 3095592H1 2/6	567
42 474800.7.dec 3015	133511		534
42 474800.7.dec 1338	/2011	42 474800.7.dec 204740111 076	522
474900 7 dec 1551	JUUI -	42 474800.7.dec 5212020111 076	501
474800 7 dec 1530	539H1 276	474800.7.dec 276583011 276	578
42 7 400 7 400 2003	256H1 277	42 474800.7.dec 9251/4711 255	490
42 7,000 7,00 5/26	861H1 276	01 - 474000 7 dec 516895901 200	
42 774007 7 200	818H1 276	07 42 474900 7 dec 5986292H1 253	
	1131H1 277	182 42 17400 7400 1966146H1 253	484
42 474800.7.dec 528		100 42 474800.7.dec 1966146H1 255	
42 4/4800.1.000	2690H1 277		
42 474800.7.dec 475	2690H1 277	93 .	

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	т.	- L1- 0	PCT/US00/25610
42 474800.7.dec 2289883H1			- 61/0300/25610
42 474800.7.dec 2424790114	253 479	42 474800.7.dec 32853841	
42 474800.7.dec 416602214	252 497	42 474000.7.dec 32853841	11 271 524
42 474800.7.dec 4373163U4	254 549	42 47 4822421	11 271 570
42 474800.7.d c 989826T1	254 508	42 474000.7.dec 1494042H	11 272 447
	276 745	42 474600.7.dec 1495315H	1 272
	276 657	42 474600.7.dec 899948R1	274 700
	276 542	7/4000./.dec 2010116U	1 07
45/0644H1	276 538	- 7/4000./.dec 30605001.	2 222
42 474000.7.dec 3715127H1	276 509	7/4000,/,dec 252200411	, 733
42 47-000.7.dec g1422739	276 625	42 474800.7.dec 48082424	
42 ATTOOUT, DEC 899948H1	274 582	42 474800.7.dec 47080914	
42 474600.7.dec 4356250H1	275 541	42 474800.7.dec 60514644	0.0
42 4740U.7.dec 4858015H1		42 474800.7.dec 96025744	232 509
42 474600.7.dec 5079276H1		42 474800.7.dec 266122714	232 514
42 474600.7.dec 1613653H1		42 474800.7.dec 4108170H1	
- 7/7000,/,DBC 3/03/401/4			232 496
7/40UU./.dec 0/E21070			232 467
" 7/40UU./.GEC 312720414	00		232 486
7/4000,/,000 0704050	35 606	42 474800.7.dec 1360090H1 42 474800.7.dec 3359828H1	232 435
42 474800.7 dec 100000000	37 609		399 674
42 474800.7.dec 192200414	42 600		404 627
44 474800.7 dec 03504446	42 588		417 605
42 474800.7.dec 02524000	183	42 474900 7 1000 91266718	426 804
44 474800.7.dec 1020cc7114	290	42 47000.7.uec g1523438	425 794
44 474800.7.dec g2011057	273	42 474000.7.UEC 91264895	427 811
42 474800.7.dec 173300014	222	42 474000.7.dec g1209924	428 787
42 474800.7 dec 35054511		42 47460H1	436 706
42 474800 7 dec 1700 (cm.)		47 4800.7.dec 4548929T1	440 746
	5 319	47 4600.7.dec g899619	442 786
42 474800 7 dog 50000		474600.7.dec g2017819	443 789
	4 409	474600.7.dec 6054166H1	
		40 474600.7.dec 5115520H1	
42 474-1460 4844124H1 217	466	- 7/4000,/.dec 50956514	400
40 1000.7.dec 4016409H1 216	362	" 7/40UU./.dec 1700400=0	10-
40 4702861H1 217	460	42 474800.7.dec 01010100	466 751
42 474000.7.dec 2523175H1 217		42 474800.7.dec 5421771110	468 783
42 474000.7.dec 5304920H1 222		44 474800.7.dec 0907455	261 438
42 47 JOOU.7. UBC 2366003H1 222	458	42 474800,7,dec 3047550114	261 576
42 474000.7.dec 3780782H1 226	535	44 474800.7.dec g834500	260 565
42 4740U./.dec 996082H1 220	335 377	42 474800.7.dec 256420014	261 562
42 474600.7.dec 822805R1 227	793	42 474800.7.dec 397979314	261 515
42 474600.7.dec 1996347R6 232		42 474800.7 dec geografia	262 534
42 474600.7.dec 3093975H1 232	637	42 474800.7.dec geaggalla	263 751
42 474000.7.dec 3133938H1 232	515	42 474800.7 dec 400700514	263 590
7/40UU./.dec 6204400114	511	42 474800.7 dec 27c2004114	262 529
42 474600.7.dec 1996347H1 222	529	42 474800.7 dec 2455000111	264 535
7/4000./.dec 286120414	496		262 504
T/7000./ GBC //2000/4414	498	74 4/4800 7 dec 050070111	263 543
"- 7/4000./.dec 2/050701/4	520	42 474800.7 dec 160000000	262 511
42 474800.7.dec 1970513H1 263	476 500	42 474800.7.dec 272752014	862 480
7/4000,/.dec 225650514	508		62 491
" 7/4000./.dec 252224714	522		62 426
42 474800.7.dec g1153120 260	516	44 474800.7 dec 01554000 5	64 502
42 474800.7.dec 615397014 200	641		61 441
42 474800,7,dec 303653344 202	552	42 47 1000.7.UEC 2496135H1 2/	54 580
42 474800.7.dec 4545077114	543	42 47 Jan 19 20 2011/87H1 2/	53 542
42 474800.7 dec 3048000111	525	42 47 1000.7.dec 3360431H1 24	53 531
42 474800.7.dec 475245014	512	42 4773156H1 27	
42 474800.7.dec 618190444	502	42 47 4000.7.dec 4379475H1 27	
42 474800.7.dec 3575044114	538	42 774000.7.dec g1551620 27	
42 474800.7.dec 251252514	562	42 474600.7.dec 2912032H1 27	
42 474800.7.dec g1330400	490	42 474800.7.dec 2632654H1 27	
42 474800.7.dec g1101105 265	700	47 474800.7.dec 3401603H1 27	
42 474800.7 dec 330650 444	F 4	12 474600.7.dec 3317001H1 27/	
		774600.7.dec 2768258H1 276	
40 g992293 260	TO4	- 7/4000./.dac agazasa	
42 47 July 752410H1 274		474800.7.dec g1200612 226	
40 45 370	140	474800.7 dec gospona	
	·64	474800.7.dec 141064114	
7,7600.7.dec 352307014	·	2 474800.7.dec 141961014 220	
	•	2 474800 7 dec 80000 220	. 478
	94	227 622805H1 227	446

				Table 2 o	ont.				
42	474800.7.dec	1571453H1	227	331	42	474800.7.dec	g1069389	248	570 ·
42	474800.7.dec	3450482H1	229	475	42	474800.7.dec	4718186H1	250	513
42	474800.7.dec	4297547H1	229	404	42	474800.7.d c	2507696H1	249	481
42	474800.7.dec	1982331H1	228	506	42	474800.7.dec	832375H1	249	338
42	474800.7.dec	2078474H1	228	493	42	474800.7.dec	3384841H1	250	477
42	474800.7.dec	4655615H1	228	494	42	474800.7.dec	3090981H1	248	514
42 42	474800.7.dec	4718836H1	228	501	42	474800.7.dec	2496134H1	247	476
42	474800.7.dec 474800.7.dec	2748984H1 2725538H1	228 228	490 477	42 42	474800.7.dec	2070351H1	251	468
42	474800.7.dec	2688345H1	228	477	42	474800.7.dec 474800.7.dec	3343389H1 2328523H1	251 251	497 502
42	474800.7.dec	5260612H1	228	452	42	474800.7.dec	1540212H1	281	501
42	474800.7.dec	2828878H1	261	526	42	474800.7.dec	1664483H1	281	493
42	474800.7.dec	3352277H1	261	513	42	474800.7.dec	g1954939	283	528
42	474800.7.dec	5438786H1	262	537	42	474800.7.dec	1652595H1	281	504
42	474800.7.dec	608875H1	260	498	42	474800.7.dec	3256087H1	282	517
42	474800.7.dec	g1687582	259	507	42	474800.7.dec	2664505H1	276	511
42	474800.7.dec	2528382H1	262	582	42	474800.7.dec	3133484H1	276	560
42	474800.7.dec	3161323H1	261	560	42	474800.7.dec	902991H1	274	443
42 42	474800.7.dec 474800.7.dec	3815719H1	262	552	42	474800.7.dec	2523971H1	274	457
42	474800.7.dec	5927826H1 2103114H1	261 261	552 524	42 42	474800.7.dec 474800.7.dec	4065211H1 5847748H1	277 277	554 555
42	474800.7.dec	5685850H1	304	457	42	474800.7.dec	3241608H1	277	555 521
42	474800.7.dec	g869962	317	521	42	474800.7.dec	1558275H1	228	438
42	474800.7.dec	g3446179	319	765	42	474800.7.dec	3945726H1	230	521
42	474800.7.dec	207985H1	323	566	42	474800.7.dec	3864787H1	224	382
42	474800.7.dec	2321546H1	323	562	42	474800.7.dec	5993349H1	230	506
42	474800.7.dec	4863479H1	333	572	42	474800.7.dec	4304036H1	229	481
42	474800.7.dec	g2540814	332	813	42	474800.7.dec	4359071H1	231	441
42	474800.7.dec	4503887H1	333	579	42	474800.7.dec	759905H1	228	443
42	474800.7.dec	g3679347	335	790	42	474800.7.dec	626533H1	232	480
42 42	474800.7.dec 474800.7.dec	4501531H1 227482H1	333 276	498 386	42 42	474800.7.dec	4359017H1	230	499
42	474800.7.dec	2864302H1	278	601	42	474800.7.dec 474800.7.dec	4160125H1 605146R1	231 232	486 ⁻ 789
42	474800.7.dec	4667387H1	280	536	42	474800.7.dec	641045H1	231	487
42	474800.7.dec	4587242H1	280	535	42	474800.7.dec	4774994H1	228	473
42	474800.7.dec	4013223H1	280	568	42	474800.7.dec	g1275341	231	726
42	474800.7.dec	4364726H1	281	536	42	474800.7.dec	5218085H1	232	489
42	474800.7.dec	5585051H1	281	508	42	474800.7.dec	1581422H1	231	414
42	474800.7.dec	g1614356	276	430	42	474800.7.dec	2875181H1	231	416
42	474800.7.dec	5292222H2	281	476	42	474800.7.dec	4108876H1	232	509
42 42	474800.7.dec 474800.7.dec	4014021H1 1996347T6	281 283	555	42	474800.7.dec	763465H1	232	458
42	474800.7.dec	2603087H1	282	749 565	42 43	474800.7.dec	1360090F1	232	712
42	474800.7.dec	2726460H1	283	533	43 43	427883.13.dec 427883.13.dec		1	585 274
42	474800.7.dec	2520565H1	282	530	43	427883.13.dec		66	600
42	474800.7.dec	g3648049	382	787	43	427883.13.dec		177	680
42	474800.7.dec	g3785151	392	789	44	018945.1.dec		148	381
42	474800.7.dec	g4621991	392	788	44	018945,1.dec		1	265
42	474800.7.dec	4548927T1	396	746	44		2494157F6	100·	578
42	474800.7.dec	2417032H1	256	496	44		5926973H1	260	527
42	474800.7.dec	2458318H1	256	473	44		5843114H1	340	556
42 42	474800.7.dec	g899701	257	366	44		2494157H1	101	374
42	474800.7.dec 474800.7.dec	g1925616 2423152H1	260 257	686 496	44 45		3332135H1	103	272
42	474800.7.dec	g1969694	257 259	496 528	45 45	353271.2.dec 353271.2.dec	g2166108 g1982479	1	268
42	474800.7.dec	g1991227	259	523	45	353271.2.dec	1330982H1	27 1	253 223
42	474800.7.dec	3984471H1	258	526	45	353271.2.dec	g656570	46	300
42	474800.7.dec	3797251H1	256	420	45	353271.2.dec	6264802H1	166	684
42	474800.7.dec	2515639H1	256	377	45	353271.2.dec	g5393964	500	874
42	474800.7.dec	1982495H1	258	434	45	353271.2.dec	g2817460	555	977
42	474800.7.dec	3384040H1	258	518	45		g3144408	555	951
42	474800.7.dec	4340308H1	262	520	45	353271.2.dec	g2884545	555	872
42	474800.7.dec	5485719H2	263	536	45	353271.2.dec	6433345H1	685	1075
42	474800.7.dec	3399655H1	249	488	46	221686.2.dec	g1496021	950	1174
42 42	474800.7.d c	4079842H1	247	504	46	221686.2.dec	2222469H1	980	1228
42	474800.7.dec 474800.7.dec	g2018263 3595893H1	248 250	489 545	46 · 46		3702201H1	4	224
42	474800.7.dec	4353689H1	250 248	466	46	221686.2.dec 221686.2.dec	2869444H1 2608081H1	5 · 5	267 239
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46		Table	2 cont.		·	PCT/US0	0/25610
46 221686.2.dec 25605	47H1 20			•			0/25010
46 22 1086.2.dec 86527	4H1 20	262 229	46	221686.2.de	c g1970171		
46 221686.2.dec 46004	87H1 1207	1471	46	221686.2.de	C 03050450	278	529
46 221000.2.dec g5395	517 1212	1658	46	221686,2,de	C 0530000	319	551
46 201000.Z.u c g3649	326 1107		46	221686.2.de	C 04300110	196	551
46 2010-2.dec 192400)3H1 120E		46	221686.2 de	04244004	200 205	564
	67 786	1285	46 46	221686.2.dec	g2986647	221	549 567
46 221686.2.dec 656089 46 221686.2.dec 430920	1H1 824	1412	46	221686.2.dec	g3134329	222	567 564
40 221686,2 dec g19046		1156	46	221686.2.dec	2192270H1	506	567
40 221686.2.dec 1330co		1068	46	221686.2.dec	3972831F8	584	948
40 221686,2,dec 133350		977	46	221686.2.dec 221686.2.dec	g1847092	601	656
46 221686,2,dec 010620		1012	46	221686.2.dec		613	903 .
46 221686,2,dec 015019		1161	46	221686.2.dec		508	785
40 221686,2,dec 2778131	⁾⁴ 748 1 1 759	1240	46	221686.2.dec	g705571	508	765
24 1000,2 dec 8016061	R1 780	997	46	221686.2.dec	g900050	543	821
46 221066.2.dec 891686H	11 780	1232	46	221686.2.dec	g2110278 3644153H1	551	941
46 221000.2.dec 6541682	H1 785	1057 835	46	221686.2.dec	3972831H1	563	845
46 221000.2.dec 2757069	H1 4	264	46	221686.2.dec	5264943H2	584	763
46 001000.2.0ec 33/1090	H1 4	191	46	221686.2.dec	91496022	583	842
46 201000.2.uec g115716	1 1	376	46 46	221686.2.dec	2276917H1	1224 1233	1649
	• -	230	46 46	221686.2.dec	92223945	448	1474
	11 3	291	46	22 1000.2.dec	g748720	489	564 816
46 221686.2.dec 2755672L	4	321	46	221686.2.dec	508694H1	492	816 564
46 221686.2.dec 45303631		281	46	22 1000.2.dec	g3871207		594
46 221686,2,dec 45303541	4 400-	1532	46		1449534H1		725
46 221686.2.dec 0977000	400	1529			93599990		1649
221686.2.dec g3085798		321	46		3212603H1		96
46 221086.2.dec 602072H1		65	40 2	21686.2.dec 1	2692989H1 524060H1	20 2	234
46 201000.2.dec g4089352	4000	i49 649	40 2	21686.2.dec c	1876029		42
46 221686.0 4 93424232	4	655	40 2	21686.2.dec 3	350967H1		37
46 001000.2.dec 9618384	4000	649	40 2	21686.2.dec 5			40
46 201000.2.UEC 9899949	1205 1	200	40 2	21686.2.dec 🚜	349946H1		343
	1217 14	470	46 2	21000.2.dec g	5368644	·	409 85.4
	1219 16	548	46 2	21000.2.dec 2	561462H1	4404	654 852
40 221686.2.dec 220150014	1484 16	***	46 22	-1000.2.dec 4	338064H1		553 199
40 221686.2.dec g3507614		48 4		1000.2.dec 65	55559H1 ·		397 397
40 221686.2.dec gassaga		54 4			100441	115 56	
40 221686.2.dec ganconzo	1536 16 1555 16		16 22		244273	20 56	
46 221000.2.dec g4619485		40	22	1686.2.dec 05		21 56	6
46 221000.2.dec g4285953	440=	40	6 22	1686.2.dec 04	2000 4 -	26 56	9
46 221086.2.dec g2670165	1465 164 1480 164	10	o 22	1686.2.dec 016	040700	37 56.	
46 221696 2 2191582H1	1 245		o 22 [.]	1686.2.dec 04/	20000		
46 221686.2.dec 5023657H1	1 526	• • • • • • • • • • • • • • • • • • • •	D 221	686.2 dec		49 602 71 591	
	949 128		221	000.2.dec g32	30774		
46 221686,2,dec g2044040	1308 145			000.2.dec g52	35938 11	188 165 190 165	
46 221686.2.dec 01531905	1313 165	6 46		686.2.dec 923 686.2.dec 194	35846 11	89 165	
46 221686.2.dec gaggious	1315 1649	9 46	:		5431H1 50		
40 221686.2 dec 04111170	1315 1649		221		1607H1 33	272	
40 221686,2,dec gs7800c	1322 1648 1328 1649		2210	586.2 dec - 634	20565 27	476	
46 221686.2.dec g3239905		, ,,	2216	86.2.dec 454	79118 12 5185H1 87	57 163	
46 221086.2.dec 3009995H1	1365 1652 1365 1654		2216	86.2.dec 361/			
46 221000.2.dec 2768117H1	75 324		2216	86.2.dec 1756			3
46 221005.2.dec g2056693	90 482	46	2216	86.2.dec 1750		398	÷
46 201000.2.dec 1914093H1	43 290	46	2216	86.2.dec 2520	533H1 7 988H1 7	260	
46 = 1000.2.uec g5664064	104 570	46 46	2216	86.2.dec 6570	842H1 999	239	
	1084 1647	46 46	2216	86.2.dec g222	9468 118		
46 221686.2.dec 1985587H1 46 221686.2.dec 2794944H1	1099 1349	46 46	2216	00.2.dec g150	1805 119		
~~ 1000.2.08C A76E00011.	1118 1379	46	22168	0.2.dec 8431(00R1 100		
40 221686.2.dec g3434999	329 635	46	22166	0.2.dec 8431(00H1 100		
40 221686.2.dec 340220014	1280 1649	46	22160	10.2.dec 13225	09T6 101		
46 221686.2.dec g2464004	227 509	46	22160	6.2.dec 28658	0.4114	312	
46 221686.2 dec 01120049	229 563	46	22169	6.2.dec 34940 6.2.dec 34219	81H1 on	141	
46 221686.2.dec 02057007	253 564	46	22168		65H1 20	225	
	564	46 -	22168			573	
*		96		6.2.dec 42002	56H1 185	456	
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Table 2 cont. 221686.2.dec g3053185 233347.7.dec 628818H1 221686.2.dec q4113264 233347.7.dec 5550441H1 221686.2.dec 1566145H1 233347.7.dec 621350H1 221686.2.dec 4146653H1 233347.7.dec 5855889H1 221686.2.dec g2038320 233347.7.dec g4331585 221686.2.dec 870643R1 233347.7.dec 1997765R6 221686.2.dec 870643H1 233347.7.dec 1997765H1 221686.2.dec 2472630H1 233347.7.dec 6117878H1 221686.2.dec 5642893H1 233347.7.dec 2241265H1 221686.2.dec g4089330 233347.7.dec g2969034 221686.2.dec g1874878 233347.7.dec 4509861H1 g760468 221686.2.dec 233347.7.dec 5075055H1 221686.2.dec 233347.7.dec q1875083 1724406H1 221686.2.dec 4115809H1 233347.7.dec 3074869H1 221686.2.dec 2190470H1 233347.7.dec g1158164 221686.2.dec g5365668 233347.7.dec 2506817T6 221686.2.dec g3665314 233347.7.dec 1683282T7 221686.2.dec g4896697 233347.7.dec 1444691H1 221686.2.dec g4682121 233347.7.dec 5201751H1 221686.2.dec g5101618 233347.7.dec 2275646H1 q3254734 221686.2.dec 233347.7.dec 2275638H1 221686.2.dec 837519H1 233347.7.dec 572758H1 221686.2.dec g2110810 233347.7.dec 1997765T6 221686.2.dec 6414447H1 233347.7.dec 2005649H1 221686.2.dec 1322509F6 233347.7.dec 2995962H1 5327256H1 221686.2.dec 233347.7.dec 4668682H1 221686.2.dec g1521860 233347.7.dec 5024735H1 221686.2.dec 1544501H1 233347.7.dec g1137365 221686.2.dec 2458467H1 233347.7.dec 4029754T6 221686.2.dec 2494070H1 233347.7.dec 2879058T6 221686.2.dec 1503112H1 233347.7.dec 782138T6 221686.2.dec g3871897 233347.7.dec 2882223T6 221686.2.dec a3085806 233347.7.dec 2048496H1 221686.2.dec g682225 233347.7.dec 2855651H1 221686.2.dec g561064 233347.7.dec 2095114H1 221686.2.dec q5707032 233347.7.dec 2882168T6 221686.2.dec 1871308H1 233347.7.dec 2132811T6 221686.2.dec 2253510H1 233347.7.dec 690448H1 221686.2.dec 3492962H1 233347.7.dec 2667831F6 233347.7.dec 5093226H1 233347.7.dec 2667777H1 233347.7.dec 6296882H1 233347.7.dec 4645421H1 233347.7.dec g1956049 g3094666 233347.7.dec 233347.7.dec 2280565H1 233347.7.dec 136644F1 233347.7.dec 4950593H1 233347.7.dec g3279211 233347.7.dec g1266199 233347.7.dec g5393561 233347.7.dec 2582086H1 233347.7.dec g2328918 233347.7.dec g3801124 3289913H1 233347.7.dec 233347.7.dec 5481292H1 233347.7.dec 6131567H1 233347.7.dec 1483352H1 233347.7.dec 3522523H1 233347.7.dec 2958439H1 233347.7.dec q4150559 233347.7.dec 5174119H1 233347.7.dec 1864389T6 233347.7.dec 573115H1 233347.7.dec 5584776H1 233347.7.dec 4110287H1 233347.7.dec 747304H1 233347.7.dec q1635141 233347.7.dec 841354H1 233347.7.dec g2027455 233347.7.dec 841354R1 3468767H1 233347.7.dec 233347.7.dec 2667831T6 233347.7.dec g1101060 233347.7.dec 3883246H1 233347.7.dec 2747911H1 233347.7.dec 1996207H1 233347.7.dec 1335579H1 233347.7.dec g660429 233347.7.dec 3280038H1 233347.7.dec 1420378H1 233347.7.dec 1394941H1 233347.7.dec g3797815 233347.7.dec 2485558H1 233347.7.d c g4094464 233347.7.dec 1441111H1 233347.7.dec g3595501 233347.7.dec 1864633H1 233347.7.dec g1211828 233347.7.dec 5289668H1 233347.7.d c g3922052 233347.7.dec 632354H1 233347.7.dec g3280797 233347.7.dec 1432029H1 233347.7.dec 053627H1 233347.7.dec 1690635H1 233347.7.dec 2573037H1

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47 233347.7.dec g3801129	Tal	ole 2 cont.	PCT/US00/25610
4/ 233347.7.dec 04091100	1214 1602	47 2200 47 -	
4/ 233347.7.dec cansent	1218 1600	47 2000 J. J. dec g10876	159 488
4/ 233347.7.dec 04242700	1218 1598	47 200047.7.dec 157051	OH1 630 045
47 233347.7.d c 04451162	1229 1602	47 20047.7.uec 221218	7H1 651
47 233347.7.dec 01110275	1230 1602 1247 1602		/H1 664
20034/./.dec 03/33000			5H1 627 637
47 233347.7.dec 1255671H1	4000	4/ 233347.7 dec 2270c4	_,
47 200047-7-Dec 3367489H1	100	47 233347.7.dec g1992c	_, 054
47 233347.7.dec g875829	1264 1532 1276 1613	47 233347.7.dec 1547400	302
47 200047.7.UBC 817888H1	1284 1439	4/ 233347.7.dec 364247	LI4 444
	1284 1597	47 233347.7.dec 2768515	H1 112 274
4/ 233347.7 dec 26004884	1284 1568	47 222247.7.dec 3292537	H1 114 257
47 233347.7 dec 01745050	1287 1573	47 200347.7.dec 3584195	H1 114 407
233347.7.dec gazagaza	1287 1597	47 200047.7.dec 5174003	H1 123 200
233347.7.dec 03744007	1290 1604	47 000047.7.UBC 552545H	1 99 249
4/ 233347.7.dec 185050070	1294 1599 1297 1550		11 101 040
47 233347.7.dec g4018301	1000	4/ 233347.7.dec. 5507404	
47 233347.7.dec 2603385H1	1297 1607 1301 1585	-0004/./.DEC 31404401	14
47 200047.7.dec 1751069H1	1302 1552	4/ 233347.7.dec 03101100	1000
	1304 1567	4/ 233347.7.dec 120ecczu	1 1007
4/ 233347.7.dec offeeten	1307 1510	47 200547.7.dec g2279662	1305 1040
4/ 233347.7 dec 01444700	1318 1603	47 200347.7.dec 2233346H	1 1395 1642 1 1396 1599
4/ 233347.7 dec 010ccom	1327 1597	47 200047.7.dec 93919171	1404 4000
4/ 233347.7.dec 0214045-	1328 1597		1412 4500
47 233347.7.dec 1211548T1	1338 1598 1343 1559		1412 4500
47 200547.7.dec 1211548R1	1343 1559 1343 1597	4/ 233347.7.dec 1630051114	1420 1529
47 200547.7.dec 1211548H1	1343 1587	4/ 233347.7.dec 02464360	1002
47 0000 1211827H1	1343 1554	47 233347.7.dec 2088964H1	1430 1602 1433 1592
4/ 233347.7.dec 93307014	1351 1603	47 200547.7.dec g879584	1002
4/ 233347.7.dec 82307074	1361 1588	47 200047-7-dec 6180192H1	115 435 116 381
4/ 233347.7.dec 205020700	361 1567	47 200547.7.dec g4530586	115 1489
4/ 233347.7.dec 205020770	362 1601	47 2000 17. UBC 2734994H1	118 366
47 200047.7.dec 2059287H1 1	362 1560 362 1601	4/ 233347.7.dec 5222000144	120 356
47 200547.7.dec 6508958H1 1	362 1601 366 1602	47 233347.7.dec 63003014	704 958
47 200547.7.dec 3886266H1 1	366 1597	*/ 233347.7.dec 4110107114	723 948
47 0000 17 Just 6508772H1 11	366 1597	47 233347.7.dec 3373756H1	739 924 735 974
4/ 233347.7.dec gagggara	366 1602	47 233347.7.dec 4816662H1	0, 4
4/ 233347.7.dec 01745050	368 1608	47 200347.7.dec 1864389H1	92 341 90 290
4/ 233347 7 dec assistante	78 1601	47 200047.7.dec g870430	92 392
4/ 233347.7.dec 227144414	82 1603		96 342
47 255547.7.dec 2596829H1 12	85 1650	4/ 233347.7 dec 1220000	97 437
47 200547.7.dec 4575094H1 12	7 270	233347.7.dec 142070014	538 731
47 20047.7.dec 3488933H1 12	7 400	47 233347.7.dec 542207014	538 727
4/ 233347.7 dec 467004 cul	625	47 233347.7.dec g2025391	536 781 538 744
4/ 233347.7.dec 21220115	406	47 233347.7.dec 705632H1	538 744 335 563
4/ 233347.7.dec 1864200Fe	783	47 233347.7.dec 2191523H1	339 651
47 233347.7.dec 4533085H1 143	584	47 100047.7.dec g1958225	361 762
47 233347.7.dec 2986556H1 143	1 1644	233347.7.dec 125000011	410 716
47 233347.7.dec g3001042 145	2 1557	233347.7.dec 2122014114	417 614
47 0000 H. J. Lee 98/0389 145	1011	233347.7.dec 782120D0	442 702
47 0000 / 16558H1 1450		233347.7.dec 4318806H1	206 698 214 478
4/ 233347.7.dec 200701711	1601	233347.7.dec g1147033	214 478 216 522
4/ 233347.7.dec c1153400	1589 4	233347.7.dec 383152H1	219 506
4/ 233347.7.dec 3409407144 14/2	1602 47		254 501
47 233347.7.dec 1429459H1 1503	1500	233347.7.dec 5262088H2	273 486
47 200547.7.dec 2130011H1 1550	1000	233347.7 dec 464000011	273 522
47 233347.7.dec 2530774H1 666	1602 47 911 47	233347.7.dec 86041014	279 534
47 233347.7.dec 136644H1 600	900	233347.7.dec 960440D4	81 533
47 233347.7 d c 49335551 690	1241 47	233347.7.dec 1683282F7 2	81 827 95 698
4/ 233347.7.dec 2077255114	955 47	200047.7.dec 2681759H1 3	95 698 12 549
159	411. 47	233347.7.dec 3950419H1 10	92 484
•	98		95 416
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				Tab	le 2 cont.		<i>"</i>		
47	233347.7.dec	g1301265	202	647	48	230631.3.dec	568183T6	1530	2106
47	233347.7.dec	782138H1	206	440	48	230631.3.dec	2611995T6	1550	2099
47	233347.7.dec	2925307H1	160	433	48	230631.3.dec	3186434H1	1585	1904
47	233347.7.dec		162	393	48	230631.3.dec	g2100758	1686	2142
47 47	233347.7.dec	2837523H2	162	401	48	230631.3.dec	g5396058	1705	2147
47 47	233347.7.dec	1520110H1	81	279	48	230631.3.dec	435524R6	1712	2026
47	233347.7.dec 233347.7.dec	643890H1 2893074H1	81 81	323	48	230631.3.dec	6314425H1	1	571
47	233347.7.dec		82	349 586	48 48	230631.3.dec	5512314F6	134	411
47	233347.7.dec	2506817H1	82 82	327	48	230631.3.dec 230631.3.dec	5512314H1 6124182H1	134	407
47	233347.7.dec	2210548H1	90	221	48	230631.3.dec	6352048H2	171 279	652 524
47	233347.7.dec	g1993764	114	386	48	230631.3.dec	5151685F6	432	780
47	233347.7.dec	4671365H1	115	382	48	230631.3.dec	5151685H1	432	697
47	233347.7.dec	5196870H1	115	305	48	230631.3.dec	g2100757	477	966
47	233347.7.dec	_ : : : : : : : : : : : : : : : : : : :	115	381	48	230631.3.dec	g1014169	477	568
47	233347.7.dec	5370648H1	111	243	48	230631.3.dec	3750444H1	494	773
47 47	233347.7.dec	2993361H1	1	316	48	230631.3.dec	5545516H1	750	948
47	233347.7.dec 233347.7.dec	3765582H1 g2011491	36	324	48	230631.3.dec	747404H1	1112	1344
47	233347.7.dec	3759276H1	108 107	403 412	48	230631.3.dec	2611995H1	1172	1415
47	233347.7.dec	2364631H1	107	339	48 48	230631.3.dec 230631.3.dec	6109396H1 5605261H1	846 965	1159
47	233347.7.dec	3660890H1	111	358	48	230631.3.dec	5101319H1	1033	1231 1301
47	233347.7.dec	2879058F6	457	839	48	230631.3.dec	568183H1	1070	1382
47	233347.7.dec	g1376379	458	816	48	230631.3.dec	568183R6	1070	1634
47	233347.7.dec	6138586H1	464	763	48	230631.3.dec	3804195H1	1086	1385
47	233347.7.dec	1930146H1	464	716	48	230631.3.dec	2759470H1	1096	1366
47	233347.7.dec	6116446H1	464	746	49	335146.1.dec	2695263F6	536	1060
47 47	233347.7.dec 233347.7.dec	2879058H1	456	781	49	335146.1.dec	g2719161	570	699
47	233347.7.dec	6138487H1 3038850H1	464 484	763 769	49	335146.1.dec	g4891403	1	447
47	233347.7.dec	5292177H2	485	769 713	49 49	335146.1.dec	g4891292	1	462
47	233347.7.dec	g828019	491	831	49	335146.1.dec 335146.1.dec	g4311647 g3887245	1	414 309
47	233347.7.dec	1850504H1	499	809	49	335146.1.dec	878550H1	247	471
47	233347.7.dec	4615174H1	506	772	49	335146.1.dec	g4266690	365	696
47	233347.7.dec	3811338H1	506	784	49	335146.1.dec	2695263H1	536	664
47	233347.7.dec	2278915H1	509	782	50	337160.1.dec	g3239640	1312	1540
47	233347.7.dec	2687896H1	511	760	50	337160.1.dec	g2080803	1319	1543
47 47	233347.7.dec	2595364H1	111	352	50	337160.1.dec	g4269757	1329	1533
48	233347.7.dec 230631.3.dec	3576583H1	104	410	50	337160.1.dec	g5633598	1271	1537
48	230631.3.dec	435524H1 435524T6	1712 1712	1808 2104	50 50	337160.1.dec	g1421937	1279	1536
48	230631.3.dec	g1927614	1714	2138	50 50	337160.1.dec 337160.1.dec	5167368H1 g752491	563 600	803
48	230631.3.dec	6410396H1	1719	2030	50 50	337160.1.dec	g1422034	609 686	905 1027
48	230631.3.dec	g4311787	1727	2143	50	337160.1.dec	5490664H1	696	990
48	230631.3.dec	g3214289	1738	2142	50	337160.1.dec	3749493T6	916	1491
48	230631.3.dec	g2567651	1772	2007	50	337160.1.dec	g3593694	1073	1447
48	230631.3.dec	g2901561	1772	2007	50	337160.1.dec	3749493H1	6	294
48 48	230631.3.dec 230631.3.dec	g5452257	1778	2147	50		g5232677	1075	1538
48	230631.3.dec	3870968H1 g2898784	1783 1802	2068	50 50		g1925336	11	448
48	230631.3.dec	g656370	1820	2007 2163	50 50	337160.1.dec	g5526629	1109	1538
48	230631.3.dec	1723586T6	1844	2102	50 50		g3596807	1110	1533
48	230631.3.dec	6556178H1	1845	2142	50		g2106751 5923530H1	105 108	561 394
48	230631.3.dec	6551620H1	1845	2159	50		4402877H1	135	389
48	230631.3.dec	1430637H1	1176	1437	50		g1164166	349	674
48	230631.3.dec	6558462H1	1845	2158	50	337160.1.dec		440	732
48	230631.3.dec	6551720H1	1845	2142	50	337160.1.dec	3457353T6	1110	1488
48	230631.3.dec	0	1866	2147	50	337160.1.dec	g752492	1196	1538
48 48	230631.3.dec	g4688016	1866	2147	50	337160.1.dec		1271	1537
48 48	230631.3.dec	g1844455	1896	2145	50		3457353F6	1	483
48	230631.3.dec 230631.3.dec	g2208492 4255443H1	1906	2144	50 50	337160.1.dec		1	247
48	230631.3.dec	2861393F6	1222 1916	1473 2145	50 51	337160.1.dec		6	343
48	230631.3.dec	2861393H1	1916	2145	51 51	346341.12.dec 346341.12.dec	909/19/	1936	2305
48	230631.3.dec	4720434H1	1922	2028	51 51	346341.12.dec		1936 1955	2242
48	230631.3.dec	294274H1	1962	2100	51	346341.12.dec		1955 2030	2218 2109
48	230631.3.dec	5445032H1	2036	2146	51	346341.12.dec		2057	2333
48	230631.3.dec	5288940H1	1407	1622	51	346341.12.dec	g1952108	2153	2448

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	Tal	ole 2 cont.	PC	T/US00/25610
51 346341.12.dec g4085	571 0404	ne 2 cont.	-	
340341.12.dec 26552	1481 0047	51 34634	11.12.dec 6096790H1	
340341.12.dec 36780	6441 0000	51 34634	1.12.dec g1983854	1892 2171
340341.12.dec d107/	712 0015	51 34634	1 12 doc 4000 4000	1914 2201
910341.12 npc /6146		51 34634	1.12.dec 4602467H1	1 269
940341.12 dec 27177	ELI4 2000	51 34634	1.12.dec 2944936H2	71 351
340341.12.dec 406123		- 1007	1.12.dec 5686792H1	84 352
949341.12 dec 407056		- 1007	1.12.dec g1985487	117 444
940041.12 dec 467440		- 1007	1.12.dec 5909568H1	155 438
51 346341.12.dec 724937			1.12.dec 2930688H2	214 506
340341.12.dec 185017	A164	U 10071	.12.dec 5447067H1	237 499
51 346341.12.dec 788800		51 346341	.12.dec 2754701H1	373 628
51 346341.12.dec 2238012	~	51 346341	.12.dec 3402751H1	511 769
51 346341.12.dec 788800		0,0041	.12.dec 1613125H1	535 715
51 346341.12.dec 1212594		51 346341	.12.dec 6161920H1	656 889
970341.12 (66 /114676)		V 100 V 1	12.dec 6162985H1	656 889
940341.12 dec 500coc		U 10041,	12.dec 2508603H1	821 1067
5 54034 1.12 dec 503450a		U 10071,	12.dec 637104H1	851 1117
340341.12 dec 2700504	LI4 0000	51 346341	12.dec 632668H1	851 1097
51 346341.12.dec 5034566			12.dec 6452454H2	1014 1468
51 346341.12.dec 2918021	114	- 10071.	12.dec 5500118H1	1149 1417
51 346341.12.dec 4548080		T 10071.	12.dec 1525276H1	1153 1384
349341.12.dec 45406541	0400		12.dec 967743H1	1165 1426
340341.12.dec 12512141		- 10041.1	12.dec 4248617H1	1166 1399
340341.12.dec 379/2001	0100	4 10041.1	2.dec 3333021H1	1165 1418
340341.12.dec 2767/06L		100 7 1 . 1	2.dec 4106103H1	1184 1443
940341.12.dec 145740cu	14 000	- 10041.1	2.dec 2135513H1	1170 1427
340341.12.dec 1/674461	11 3201 3374		2.dec 5430803H1	1181 1421
940341.12.0ec 47140001	14	- 10071,1	2.dec 3287061H1	1212 1319
340341.12.dec 452226011			2.dec 5430764H1	1230 1489
51 346341.12.dec 1868560H	0403		2.dec 5953804H1	244 1569
340341.12.dec 187007011		51 346341 12	2.dec 3002616H1	251 1494
51 346341.12.dec 446846H1	0.107	- 10071.12	2.dec 1793327H1 1	262 1512
51 346341.12.dec g3134178		- 10071.12	2.dec 5505305H1 1	268 1486
540541.12.dec n5425020	3254 3631		dec 2159582H1 1	273 1431
340341.12.dec m2836424	3259 3718	- 10071.12	dec g2238358	317 1591
340341.12.dec 2120270L	3262 3716		dec 6456333H1 13	365 2013
340341.12.dec 2404565U4		J.0071.12	dec 3161945H1 13	385 1661
340341.12.dec 188001EU4		- 100-11,12,		1607
340341.12.dec n324620E	222	51 346341 12	dec 890596H1 14	159 1730
540341.12.dec 65503/12	3284 3716	51 346341 12	dec 899800H1 14	75 1752
340341.12.dec n3101//En	3308 3717	T 10071.12.	dec g4307656 14	80 1829
070341.12.0ec n5636360	3314 3717	- 10071.12.	dec 1231448H1 14	94 1734
21 040041.12.dec 04738222	3316 3716	00 + 1, 12,1	dec g3336553 14	
340341.12.dec 3218761U4	3319 3716	51 346341 12	dec 2991495H1 15	
- 070341,12,0ec 53/0743U4	3328 3628	51 346341 12	dec 4066993H1 15	17 1788
- 070041.12.0ec n3791515	3337 3539	51 346341 12 6	dec 4801741H1 15: dec 1992208H1 15:	
- 040341.12.dec 5187100U4	3354 3716		lec 4958003H1 156	28 1821
540341.12.dec 464224704	3378 3600	51 346341 12 4	ec 3153981H1 156	7 1824
540341.12.dec 154704404	3379 3646	51 346341 12 d	00 200000=+++	' 3 1845
540541.12.dec d900500	3381 3590	51 346341 12 d	00 640000000	'8 1828
340041.12.dec d2620224	3383 3726	51 346341 12 d	ec 3758378H1 158	1 2202
540341.12.dec 02069144	3388 3716	51 346341 12 d	ec 3/583/8H1 160 ec 3405995H1 158	5 1787
340341.12.dec d3837364	3392 3716	51 346341 12 0	200000	7 1825
340341.12.dec n3110249	3408 3823	51 346341 12 de	30 3FC40004	7 1885
340341.12.dec n2010076	3406 3825	51 346341 12 46	NO 200000000	
340341.12.dec a767800	3413 3717	51 346341 12 40	© 3000290H1 1602 © 1892929H1 1604	2 1885
340341.12.dec 0608251	3426 3718	51 346341 12 do	o FOODOATIA	1867
340341.12.dec d2605526	3451 3718	346341.12.de	0 -40004	
340341.12.dec g3756410	3478 3716	1 346341 12 do	- 6400	1874
340341.12.dec a796776	3517 3716	~	0.61000=	1934
340341.12.dec d301017c	3518 3729	T 100 T 1. 12.UE	. 225000	1890
91 040041.12.dec 02320740	3518 3716	- 10071.12.UE		1870
340341.12.dec 3800/63H4	3522 3716	T. 100 T 1. 12.0E	C 3340411U4 4646	
340341.12.dec d2768086	3523 3612 5	- 10071.12.080	379212744 4444	1925
340341.12.dec 062801U1	3537 3716 ₅		: 424009AU4 4000	1986
540341.12.dec 640007014	1872 2137 5		121814604	1872
91 949341.12 dec 22144661.	1880 2442 5		5079596U4 4666	1853
	1881 2132 5	· 040341.12.080	3850521114	1871
	•	346341.12.dec	6141523H1 1630	1890
	100			

				Tabl	e 2 cont.		r	C 170500	J/2501U
51	346341.12.ded		1630	1914	54	245000.6.dec	1355155H1	23	281
51	346341.12.dec		1630	1872	54 ·	245000.6.dec		40	612
51	346341.12.dec		1634	1874	54	245000.6.dec	3189465H1		394
51 51	346341.12.dec		1634	1884	54	245000.6.dec			314
51 51	346341.12.dec		1644	1895	54	245000.6.dec			301
51	346341.12.dec 346341.12.dec		1644	1822	54	245000.6.dec			292
51	346341.12.dec	. 454130/H1	1648 1665	1916 1863	54 54	245000.6.dec		50	263
51	346341.12.dec		1719	1878	54 54	245000.6.dec 245000.6.dec		50	123
51	346341.12.dec		1746	1916	54	245000.6.dec	3106926H1 2099640H1	27 50	305
51	346341.12.dec	4252513H1	1735	1896	54	245000.6.dec	3670326H1	50 57	278 341
51	346341.12.dec	1998609H1	1736	1872	54	245000.6.dec	853792H1	1	245
51	346341.12.ded	4543814H1	1754	1860	54	245000.6.dec	2512562H1	12	348
51	346341.12.dec	4111257H1	1759	2011	54	245000.6.dec	2850207H1	23	319
51 51	346341.12.dec	3239986H1	1847	2096	54	245000.6.dec	3452542H1	22	260
51	346341.12.dec 346341.12.dec		1872	2078	54	245000.6.dec	1355155F1	23	460
52	428745.2.dec	g2820616	1872 193	2102 532	54	245000.6.dec	4539474H1	59	331
52	428745.2.dec	g2881239	230	653	54 54	245000.6.dec 245000.6.dec	2563931H1 3085485H1	57 50	330
52	428745.2.dec	g1063734	241	491	. 54	245000.6.dec	3926193H1	58 59	315 342
52	428745.2.dec	g954024	275	581	54	245000.6.dec	2460549H1	60	309
52	428745.2.dec	6490302H1	322	915	54	245000.6.dec	3037582H1	59	245
52	428745.2.dec	724811R6	436	916	54 ⁻	245000.6.dec	g1377709	58	403
52	428745.2.dec	2562118H1	516	777	54	245000.6.dec	3254001H1	60	321
52 52	428745.2.dec	5463505H1	669	858	54	245000.6.dec	2691264H1	60	318
52 52	428745.2.dec 428745.2.dec	918149H1 g1991232	1	229	54	245000.6.dec	4669568H1	60	327
52	428745.2.dec	g2028636	22 116	303 368	54 54	245000.6.dec	2514525H1	62	392
52	428745.2.dec	5086711H1	124	374	54 54	245000.6.dec 245000.6.dec	2209841H1 851811H1	62	310
52	428745.2.dec	5086711F6	124	666	54	245000.6.dec	4599225H1	62 61	327 349
52	428745.2.dec	g1578109	138	560	54	245000.6.dec	1286024H1	66	318
52	428745.2.dec	g1812416	142	598	54	245000.6.dec	5075587H1	67	337
52	428745.2.dec	1713552H1	181	426	54	245000.6.dec	2515049H1	67	372
52 52	428745.2.dec	g2881535	186	328	54	245000.6.dec	5530160H1	67	335
52 52	428745.2.dec 428745.2.dec	3630275H1 4838210H1	704	849	54	245000.6.dec	g1088112	73	355
52	428745.2.dec	5078612H1	706 726	832 983	54 54	245000.6.dec	2098876H1	75	338
52	428745.2.dec	4130312H1	753	1014	54 54	245000.6.dec	1227790H1	76	349
52	428745.2.dec	2768676H1	767	1006	54	245000.6.dec 245000.6.dec	763189H1 979662H1	77 79	298
52	428745.2.dec	3448885H1	780	1025	54	245000.6.dec	2913785H1	79 81	372 346
52	428745.2.dec	4381065H1	805	1060	54	245000.6.dec	2913737H1	81	338
52	428745.2.dec	1294626T6	929	1385	54	245000.6.dec	3686742H1	82	375
52 52	428745.2.dec	1294626H1	936	1146	54	245000.6.dec	4202894H1	98	208
52 52	428745.2.dec 428745.2.dec	1294626F6	936	1423	54	245000.6.dec	2735065H1	102	318
52		1292590H1 5099834H1	936 963	1140	. 54	245000.6.dec		102	367
52		2729359H1	1041	1141 1228	54 54	245000.6.dec 245000.6.dec	2314724H1	105	333
52		2792430H1	1075	1382	54	245000.6.dec	2912582H1 139138H1	124	391
52	428745.2.dec	1888850H1	1111	1387	54	245000.6.dec	g1947635	126 124	262 336
52	428745.2.dec	6547071H1	1155	1391	54	245000.6.dec	1795035H1	137	423
53	444839.17.dec		1	507	54	245000.6.dec	5882992H1	149	419
53	444839.17.dec		221	428	54	245000.6.dec	2438787H1	148	333
53 53	444839.17.dec		310	585	54	245000.6.dec	5888569H1	149	330
53	444839.17.dec 444839.17.dec	5018181R8	552	908	54	245000.6.dec	5889878H1	150	404
53	444839.17.dec	61121716 611217H1	614 626	805	54 54	245000.6.dec	2439472H1	150	392
53	444839.17.dec	61121786	626	855 855	54 54	245000.6.dec 245000.6.dec	280152H1	624	950
54	245000.6.dec		383	640	54 54	245000.6.dec	2311893H1 983601T1	663	902
54		1929331H1	519	751	54	245000.6.dec	983601H1	685 685	1156
54	245000.6.dec	1751338H1	521	722	54	245000.6.dec	346874T6	715	971 1156
54	245000.6.dec	2515886H1	424	739	54	245000.6.dec	280952T6	747	1126
54		2326214H1	534	780	54	245000.6.dec	694271H1	159	408
54 54		280952R6	517	959	54	245000.6.dec	2413665H1	159	379
54 54		2658307H1	579	822	54	245000.6.dec	g1087215	184	452
54 54		3107867H1	27	299	54		g1799013	185	648
5 4		3114192H1 3210893H1	32 37	298	54 54	245000.6.dec	4050305H1	196	492
54		860506H1	40 ·	105 312	54 54	245000.6.dec 245000.6.dec	4052675H1	196	463
					1/11	290.0.0000	3534417H1	904	1031

	54 245000 6 dec			T	able 2	cont.		PCT/	US00/25610
	0000.0.00		11 77						
	0000.0.000	g3923886	80			55	428362.36.dec g1493	078	129 321
	54 245000.6.dec	3841719H	1 22			55	720002.30.0ec 01721/	SO2	100
	245000.6.dec	38201104	1 22			55	720302.30.dec a1492	799	
	4 245000.6.d c	4144664H	1 29			55	740004.30.dec 408941	IELI4	
	4 245000.6.dec	3689241H	1 30			55	*40304.36.dec 227538	11 LI 1	11 290
	4 245000.6.dec	24137/20	1 30			55	420302.36.dec g49297	'42	14 258
	4 245000.6.dec	5072965H2	2 37			55	420302.36.dec 340071	2111	16 908
	4 245000.6.dec	13427541	754			55	428362.36.dec g19574		16 215
	4 245000.6.dec	384005501	740			55	428362.36.dec 142317		16 419
5	5 428362.36.dec	323045611			5	55	428362.36.dec 489590	^1 • •	17 222
5	9 420302.36.dec	7468∩⊿⊔₁	00	202		55	428362.36.dec 546870	1 4 4	20 326
5	9 440302.36 dec	415887044	26	250		55	428362.36.dec g43293		20 267
5	* *40002.36.dec	8715000		294		55	428362.36.dec 1823626	~~~	242 670
55	440302.36.dec	0/100000	490	896		55	428362.36.dec g163528		255 858
55	* 740302.36.dac	04110000	500	906		55	428362.36.dec 4746558		96 683
55	428362.36.dec	02555707	502	832		55	428362.36.dec g217853		25 564
55	428362.36.dec	92333/3/ 9888606	529	900	:	55	428362 36 dog 407024	3	33 539
55	428362.36.dec	314106040	531	845		55	428362.36.dec 4973919	H1 3	35 609
55	428362.36.dec	03340000	557	805		55	428362.36.dec g215951	1 4	6 394
55	428362.36.dec	93240896	569	904		55	428362.36.dec 1998274	H1 4	B 119
55	428362.36.dec	91501350	578	879		55	428362.36.dec 2413133	H1 4(
55	428362.36.dec	94148815	803	890		55	428362.36.dec 833838H	1 52	335
55	428362.36.dec	2100865H1	20	270		55	428362.36.dec 14677591	H1 48	
55	428362.36.dec	3136845H1	22	280		5	428362.36.dec 61112411	H1 52	
55	428362.36.dec 3	0016233F6	23	517		5	428362.36.dec 835269H	1 52	
55	428362.36.dec 3	0016233H1	23	327	5.		428362.36.dec 45084031	1 1 38	
55	428362.36.dec 6	7420559H1	26	526	5	_	428362.36.dec 53090281	11 40	
55	428362.36.dec 1	949076H1	1	166	5		428362.36.dec 4352115H	11 42	
55	428362.36.dec 1	82957H1	19	81	55		428362.36.dec 2661812H	i1 45	314
55	428362.36.dec g	4074435	665	902	55		428362.36.dec 3616233T	6 34	
55	428362.36.dec g	1477032	685	905	55	_	428362.36.dec 1470246H	1 36	5 553
55	428362.36.dec g	1493079	687	908	55	•	420302.36.dec 2588112E	6 07	
55	428362.36.dec g	4330971	690	901	55	•	428362.36.dec 2588112H	1 270	
55	428362.36.dec 4:	365938H1	695	901	55 55		**************************************	373	
55	428362.36.dec g4	4510083	736	901	55	•	+20362.36.dec 02987835	400	
55	428362.36.dec 64	100671H1	757	904	55 55	•	120302.36.dec 455065gT	410	
55	428362.36.dec g2	782804	766	906		-	20362.36.dec 475983H1	100	
55	428362.36.dec 17	'40312H1	767	907	55 55	٠,	20302.36.dec 4337255H.	104	
55	720302.36.dec a1	920531	784	907		4	20362.36.dec 407178941	1 404	398
55	420302.36.dec a3	429060	793	908	55 55	4	20302.36.dec 6125624H1	155	627
55	428362.36.dec g3	093231	586	901	55	4	20302.36.dec 460302H1	210	473
55	428362.36.dec g5	035307	589	896	55 55	4	28362.36.dec 5853923H1	160	473 426
55 55	420302.36.dec a2	341502	642	902	55 55		20302.36.dec 4914457H1	174	
55	420302.36.dec a1	721060	643	908	55 55	4,	<0302.36.dec 4871852H1	1	417 278
55	420302.36.dec a21	177837	004	890	55	44	20302.36.dec 551710844	6	
55	*20302.36.dec 254	11141TE		888	55 55	44	20362.36.dec 4420770H4	8	280
55	*40302.36.dec 557	7กธวอม∢		682	55	42	0302.36.dec 5043882U4	11	268
55 55	420302.36.dec 348	39030114		292	55	42	0302.36.dec 6119524U1	12	278
55	720302.35.dec 777	'420 11 .	'	169	55	42	8352.36.dec 711540H1	65	622
55	*20302.36.dec 616	811201		526	55	42	0302.36.dec 5482010L1	68	340
55	420302.36.dec 586	5532LI4 4		313	55	42	0302.36.dec 716182H1	69	347
	*20002.36.dec 335	444641 4		303	55	42	8362.36.dec 4922936U1	71	281
55 55	420302.36.dec 323	454444 ,			55	42	8362.36.dec 323486741	71	339
55 55	420302.36.dec a15	N1340 -		270	55	42	0302.36.dec 4981082L1	75	315
55 55	420302.36.dec 257	111144 /		16	55	42	5362.36.dec 5020845U4		356
-	**COOO2.30.dec a24*	27217		67 01	55	420	3302.36.dec 2490005H4	97	333
-	420302.36.dec 5292	1749H1 4		01 05	55	420	352.36.dec 132055541	108	346
•	420302.36.dec a469	27087 A		95 24	55	428	3362.36.dec 4550658H1	52 50	300
33	+20302.36.dec a536	1720	~	01	55	428	362.36.dec 4798082H1	52	29 6
33 4	+<0302.36.dec a152	6406	~	01	55	428	362.36.dec g2161592	53	287
33 4	+20302.36.dec 0452	4022		07	56	480	710.12.dec g5235305	53	489
55 4	20302.36.dec a293	3287	78 90		56	480	710.12.dec g1557590	4035	4473
JJ 4	28362.36.dec a291	2603	79 90		56	480	710.12.dec 880717T1	4036	4470
55 4	28362.36.dec 0440	A			56	480	710.12.dec 880717R1	4038	4430
JJ 4	28362.36.dec 8715				56	480	710 12 dos 80071/H1	4038	4330
33 4	28362.36.dec 87150				56	480	710.12.dec 880717H1	4038	4275
	28362.36.dec 87150	107-		6	56	480	710.12.dec g4084804	4044	4473
	28362.36.dec g2537		9 . 86		56	480	710.12.dec 4371905H1	4050	4314
•	g2537	776 11	6 61	0		4807	710.12.dec 4506862H1	4057	4317
					,	10 0/	10.12.dec g2541764	4059	4474
				102	•				

			iabi	e 2 cont.			
56	480710.12.dec g5367783	4060	4470	56	480710.12.dec 1215902H1	3570	3798
56	480710.12.dec g678587	4067	4441	56	480710.12.dec 1657958H1	3571	3779
56	480710.12.dec g4607015	4071	4475	56	480710.12.dec g2229755	3550	4000
56	480710.12.dec g4535236	4071	4470	56	480710.12.dec 310435H1	3564	3761
56	480710.12.dec 2900362H1	4081	4354	56	480710.12.dec 4885652H1	3545	3749
56	480710.12.dec g918370	4081	4391	56	480710.12.dec 2874149H1	3715	3988
56	480710.12.dec 3836936H1	4091	4374	56	480710.12.dec 3444009H1	3729	3987
56	480710.12.dec g1692010	4092	4484	56	480710.12.dec 782063R1	3906	4445
56	480710.12.dec 1334882H1	4119	4351	56	480710.12.dec g5636497	3210	3491
56	,480710.12.dec 4861120H1	4119	4386	56	480710.12.dec 4740585H2	3218	3486
56 56	480710.12.dec 1351565F1	4125	4470	. 56	480710.12.dec 3453705H1	3219	3337
56 56	480710.12.dec 1351565H1	4125	4376	56	480710.12.dec 4350876H1	3225	3357
56	480710.12.dec 1351565F6 480710.12.dec 1377778T6	4125	4474	56 56	480710.12.dec g2059287	· 3177	3489
56	480710.12.dec g3108603	4126 4131	4441	56 56	480710.12.dec g727073	3173	3464
56	480710.12.dec 2271108H1	4132	4475 4395	56 56	480710.12.dec 554290R6	3343	3839
56	480710.12.dec 2271114H1	4132	4390	56	480710.12.dec 554290H1 480710.12.dec g2775291	3343	3563
56	480710.12.dec 4954215H1	4134	4389	56	480710.12.dec 92775291 480710.12.dec 5298647H1	3346 3350	3486 3594
56	480710.12.dec 5599026H1	4134	4378	56	480710.12.dec 3172587H1	3357	3631
56	480710.12.dec g4299288	4141	4445	56	480710.12.dec 2041156H1	3360	3625
56	480710.12.dec g813687	4143	4480	56	480710.12.dec 2766832F6	3378	3780
56	480710.12.dec g4524478	4154	4470	56	480710.12.dec 2766840H1	3378	3604
56	480710.12.dec g1266072	4169	4470	56	480710.12.dec 3270815H1	3707	3959
56	480710.12.dec g768815	4173	4472	56	480710.12.dec 5189122H1	3714	3932
56	480710.12.dec g958680	4174	4438	56	480710.12.dec 3072628H1	3285	3577
56	480710.12.dec 5104025H1	4188	4445	56	480710.12.dec 4700004H1	1600	1879
56	480710.12.dec g657133	4194	4473	56	480710.12.dec 1711787H1	1664	1877
56	480710.12.dec g715881	4195	4474	56	480710.12.dec g2031416	1698	1952
56 56	480710.12.dec g1748430	4200	4471	56	480710.12.dec 6485042H1	1723	2292
56	480710.12.dec g2556567 480710.12.dec g656966	4213	4478	56 56	480710.12.dec 4989752H1	1726	1847
56	480710.12.dec 4199390H1	4214 4230	4470 4472	56 56	480710.12.dec 4989774H1	1725	1963
56	480710.12.dec 413333011	4260	4405	56 56	480710.12.dec g1087373	1742	2115
56	480710.12.dec g1025062	4265	4433	56	480710.12.dec 130838R6 480710.12.dec 130838H1	1747 1746	2258 1928
56	480710.12.dec 598848H1	4270	4470	56	480710.12.dec g1271319	1768	2109
56	480710.12.dec 4825131H1	3886	4138	56	480710.12.dec 3291281H1	1802	1927
56	480710.12.dec 2361847T6	3888	4433	56	480710.12.dec 3692084H1	1821	2099
56	480710.12.dec 374196H1	3488	3728	56	480710.12.dec 3400646H1	1877	2103
56	480710.12.dec 2462560H1	3801	4047	56	480710.12.dec g1471362	1885	2122
56	480710.12.dec 4590905H1	3787	4017	56	480710.12.dec 3522564H1	1902	2212
56	480710.12.dec g2264982	4237	4473	56	480710.12.dec g5394524	3039	3486
56	480710.12.dec g2266184	4243	4467	56	480710.12.dec g5394523	3040	3486
56 56	480710.12.dec g5638968	4244	4470	56	480710.12.dec g3431663	3042	3494
56 56	480710.12.dec g5594372	4247	4470	56	480710.12.dec 3801488H1	3255	3509
56	480710.12.dec g1443422 480710.12.dec g761095	4250	4470	56 56	480710.12.dec 2798884H1	3265	3509
56	480710.12.dec g701095	4252 4257	4446 4470	56 56	480710.12.dec g1636310	3271	3454
56	480710.12.dec 5436689H1	3403	3598	56 56	480710.12.dec g2694542 480710.12.dec g1358669	3040 3058	3487
56	480710.12.dec 5487134H1	3431	3682	56	480710.12.dec g1338669	4297	3486 4441
56	480710.12.dec 5284219H1	3504	3657	56	480710.12.dec g1748444	4305	4471
56	480710.12.dec 2099792H1	3541	3699	56	480710.12.dec g5438233	4383	4474
56	480710.12.dec g922955	3308	3487	56	480710.12.dec 5064281H1	1910	2138
56	480710.12.dec 3614786H1	3316	3614	56	480710.12.dec g1690244	1919	2165
56	480710.12.dec 6389577H1	3321	3566	56	480710.12.dec 4539338H1	1945	2181
56	480710.12.dec g1920533	3321	3486	56	480710.12.dec 4540969H1	1945	2196
56	480710.12.dec g958726	3616	3861	56	480710.12.dec 3377229H1	2045	2296
56	480710.12.dec 5495496R6	3618	3943	56	480710.12.dec 2657748F6	2113	2548
56	480710.12.dec 884462H1	3629	3864	56	480710.12.dec 2657748H1	2113	2327
56 56	480710.12.dec g715880	3615	3900	56	480710.12.dec 4302139H1	2278	2555
56 56	480710.12.dec 5037750H1	3632	3867	56	480710.12.dec 190789R6	2288	2700
56 56	480710.12.dec 3478382H1	3651	3845	56 56	480710.12.dec 190789H1	2289	2455
56	480710.12.dec 390082H1 480710.12.dec 1845386R6	3655	3933	56 56	480710.12.dec 2971888H2	2297	2573
56	480710.12.dec 1645386H6 480710.12.dec g4149091	3669 3300	4009	56 56	480710.12.dec 3050381H1	2321	2600
56	480710.12.dec g5233011	3302	3464 3486	56 56	480710.12.dec 1363666F6	2341	2843
56	480710.12.dec 3220964H1	3303	3 4 66 3633	56 56	480710.12.dec 1363666H1	2341	2550
56	480710.12.dec g919138	3315	3635	56 56	480710.12.dec 2379137H1 480710.12.dec 3254033H1	2373	2594 2628
56	480710.12.dec 6552777H1	3570	4095	56	480710.12.dec 5056123H1	2390 2399	2628 2673
				-		2333	2013

	E0					Ta	ble 2	COnt		;	PCT/US	S00/2	5610
	56	480710.12	dec g1968	3577	2420			_			•		
	56 56	480/10.12	.dec 34507	7Ř3H1	2443			56		2732286	L1 00		
	56 56	400/10.12	dec 42738	ини	2501	,		56	7007 TO. 12. Gec	4657 <i>AA</i> U	4	06	4120
	56 56	460/10.12	.dec 43588	6641	2568			56	400/10.12.dec	1840000	Te oo	45	4176
	56 50	480/10.12	.dec 65498	81 LJ 4	2604			56	7007 10.12.dec	2722261	Te aa	48	4432
	56 56	400/10.12	.dec 14445	4441	2637	3093		56	7007 10.12.dec	27223641	مُم له	-	4443
	56 50	460710.12.	dec a1358	400	2736	2908		56	7007 10.12.dec (64R2171L	4		293
	56 50	460710,12,	dec 14367/	DRE4	2737	3179		56	4007 10.12.dec (72210055			615
	56 50	460/10.12.	dec 14367/	18LI4	2737	3258		56	7007 10.12.dec (ねろうフロウク		_	491
	56 56	400/10.12.	dec 143842	25114	2737	3006		56	700/10.12.dec d	11985224			695
	56 56	400 710.12.0	dec 162763	11 Ec	2757	2979		56	480710.12.dec	1677757L			601
	56 56	400/10,12,0	dec 162762	14114	2757	3064		56	4007 10.12.dec 3	5925151	4 400		571
	56 56	7007 10.12.0	JBC 197401	LI4	2785	2951		56	400/10.12.dec 3	3221625	6 400	-	742
	56 56	400/10.12.0	1ec 283031	2014	2798	2977		56	7007 TO. 12.dec 3	スクク16つい	1 400	_	987
	56 56	7007 10. 12.0	18C 013838	16	2839	3035		56	4007 IU. 12.dec 3	281467W	1		'51
	56 56	7007 10, 12,0	ie c 431000	2 LI∢	2847	3174		56	480710.12.dec 3	281450L			'59
	56 56	7007 IU. 12.d	ec 494817	77	2899	3123		56	400/10.12.dec 3	R24001U+		_	56
	56 56	7007 10.12.d	ec 207065	7LJ 4	2904	3448		56	4007 TU. 12.dec 5	74REELI1			63
	56 56	7007 IU. 12.d	8 C 7516691	41	2923	3179		56	4007 TO. 12.dec 41	18614544	595		23
	56 56	400/10.12.de	ec 1911 <i>4</i> 71	I E C	2925	3156 3484		56	400/10.12.dec 33	373301₽4	040		131
	30	400/10.12.de	BC 1911471	44	2925			56	7007 10.12.dec 29	168519L14	040		100
	-	4007 10.12.de	9C 4254N47	'LI4 .	2933	3200 3208	_	6	4007 10.12.dec 63	789กคม •	_		257
		400/10.12.de	C 2234211	TC .		3468		6	4007 TU. 12.dec 49	481706			160
		4007 IU. 12.0E	C 494817T	· .		3449		6	4007 TU. 12.dec 49	481707	1099		199
5		+007 10.12.de	C 2797506	Ec .				6	4007 10.12.dec 41	5823044	1099	13	116
50	•	1007 10.12.de	C 3175668	⊔ 4 ,		3425 3316		6	400/10.12.dec 52	1804144	1179		26
50	-	юи/ IU. 12.de	C 0221800			3464	5		4007 TO. 12.dec of	40634B	1363	16	
56	~ ~	10.12.de	C 01030551		`	3488	5		700/10.12.dec d1/	147767	1408	18	
56	U 4	10.12.de	C 01690120			3486 ·	5		400/10.12.dec 376	33147W4	1409	18	
56	_	007 10.12.de	C 554290TA	• •	- ·	1429	50	_	700/10.12.dec 236	184706	1435	16	
56	- 4	ου/ 10.12.deα	27668222	-		425	56	•	4007 10.12.dec 236	1847U+	1440	172	
56	•	007 10.12.dec	: 2654374L	14 4		295	56	•	7007 10.12.dec 638	2240114	1440 1458	168	
56	_	007 10.12.dec	: 1627631⊤	·e	'	429	56		^{του/} 10.12.dec 278	ハクオフレィ	1509	153	
56		907 10.12.dec	: 0445110 <i>a</i>			471	56		700/ IU. IZ.Gec 533	りりろうしょ	1541	174	
56		207 10.12.dec	: 2415120L	4 ^-		029	56 50		134 tour 10.12.dec	516144	1579	180	
56	70	707 IV. 12.Gec	241468AL	1 0-		012	56 50	_	1007 IV. 12.0ec ngo	2056	4297	181	
56		30710.12.dec	1545383H	1 ^		584	56 56	-	100/10.12.dec n75	1017	3166	447	
56		0710.12.dec	4043654H	1 34		769	56 56	_	1007 10.12.dec a380	11141	3094	347	
56	48	0710.12.dec	1845386H	1 36		952	56 56	_	207 10.12.dec a530	5425	3100	349; 348	2
56	48	0710.12.dec	9761231	36		74	56 56	7	907 10.12.dec 5297	700AU+	3106	3357	
56	48	0710.12.dec	9761199	36		104	56	-	907 10.12.dec 5296	9124	3106	3336	
56	48	0710.12.dec	g1747939	36		88	56	_	00/ 10.12.dec a567	1970		3488	
56	480	0710.12.dec	91/47953	368	35 37		56	_	00/10.12.dec d323	1367	3139	3493	
56	480	0710.12.dec	343669H1	370			56		907 10.12.dec g261	9560	04.40	3507	
56	480	710.12.dec (95446447	400)8 44°		56	71	20/10.12.dec 4117	R14LI4	_	3428	
56	480	710.12.dec c	34084680	401	3 44		56	~	0/10.12.dec 4114	106114	04	3420	
56	480	710.12.dec 3	4395415	401	4 447		56	70	''' '''. 12.0ec a110,	1822		3486	
56	480	710.12.dec 5	05047414	373		51	56	40	0710.12.dec g1383	3757		3486	
56	480	710.12.dec g	15007EA	374	4 401	0	56	40	0710.12.dec g3230	398		3486	
56	480	710.12.dec 6	305750114	377			56	48	0710.12.dec g4222	799	3055	3486	
56	480	710.12.dec 6	560125U4	377	1 433	3	56	48	0710.12.dec g1994	805		3486	
56	4807	710.12.dec 5	71050540	389	444	2	56	48	0710.12.dec g3278	265		490	
56	4807	10.12.dec 1	70527004	3897			56	48	0710.12.dec g4076		3079 з	485	
56	7007	10.12.dec 17	79597906	3842			56	480	0710.12.dec 13636	66T6 ;		439	
56	4807	10.12.dec 62	2800216U6	3842	,		56	480	0710.12.dec g2743 0710.12.dec g7611	581 ;		489	
56	4807	10.12.dec ge	38062N	3845			56	480	710.12.dec g/6112		4287 4.	463	
56	4807	10.12.dec 19	MA608U4	3859			56	480	710.12.dec 49405	37H1 4		445	
56	400/	10.12.dec a1	320173	3807			56	480	710.12.dec 598928 710.12.dec 332216	3H1 4	270 4	383	
56	4007	10.12.dec 54	871 <i>4</i> 2U+	3819			56	480	710.12.dec 332216	216 3		123	
56	4007	10.12.dec 49	05295H4	3590			56	480	710.12.dec 199531 710.12.dec 184538		873 44	131	
56	TOU /	14.12.dec 18.	40022De	3591	3852		56	480	710 12 dec 640-	616 3	874 44	133	
-	4007	10.12.dec a10	979716	3613	4077		56	480	710.12.dec 640771	4H1 3		28	
	4007	9.12.dec a16	725256	3613	3912		56	480	710.12.dec 479371	8H1 3.		34	
	40 07 1	U.12.dec 190	35310De	3616	3900			4807	710.12.dec 243320		22		
-	40071	U.12.dec 190	35310LI4	3501	3878			4807	710.12.dec 308271 710.12.dec 358955		29		
• • • •	40U/ J	V.12.dec 558	MUEULIA	3501	3760			4807	10.12.dec 358955	5H1 23	3 28		
56 . 4	48071	0.12.dec 782	063H1	3504	3732		_	4807	10.12.dec 3613921		3 25		
				3906	4120		66	4807	10.12.dec 3147763	H1 28	3 27		
						104			314//63	H1 30	289	9	

			Labic	L Cont.			
56	480710.12.dec 2722361F6	34	544	57	234137.10.dec 6407056H1	1391	1562
56	480710.12.dec 2797596H1	2948	3197	57	234137.10.dec g2964089	1397	1563
56	480710.12.dec 3729452H1	3016	3311	57	234137.10.dec g649064	1411	1567
56	480710.12.dec 4986129H1	3018	3298	57	234137.10.dec g1264136		
56	480710.12.dec 5451222H1	3027	3288	57	234137.10.dec g1264136	1380	1568
56					234137.10.dec g674420	1382	1484
	480710.12.dec 5451783H1	3026	3227	57	234137.10.dec 2110777T6	1054	1541
56	480710.12.dec g3736306	3037	3490	57	234137.10.dec 535718H1	1063	1274
56	480710.12.dec 2797596T6	3038	3447	57	234137.10.dec 1440773R1	1072	1560
57	234137.10.dec g5636768	1332	1560	57	234137.10.dec 3813579H1	855	1143
57	234137.10.dec g3873139	1336	1569	57	234137.10.dec 4937088H1	870	1121
57	234137.10.dec g830941	1337	1570	57	234137.10.dec 1993760H1	154	367
57	234137.10.dec g678033	1328	1560	57	234137.10.dec 3506477H1	154	440
57	234137.10.dec g3764879	1332	1565	57 57			
5 <i>7</i>					234137.10.dec g991858	160	319
	234137.10.dec 567963H1	724	978	57	234137.10.dec g4152671	207	561
57	234137.10.dec 643036H1	753	991	57	234137.10.dec g4737226	1154	1560
57	234137.10.dec g3434330	1204	1565	57	234137.10.dec 1516507T6	1140	1526
57	234137.10.dec g4568341	1206	1561	57	234137.10.dec g4224234	1157	1567
57	234137.10.dec g4394560	1219	1560	57	234137.10.dec 1255603T6	1172	1524
57	234137.10.dec 3150617H1	717	988	57	234137.10.dec g3096244	1181	1473
57	234137.10.dec 1440773T6	1141	1520	57	234137.10.dec 3424013H1	1000	1268
57	234137.10.dec g2138832	1142	1563	57	234137.10.dec 1742695H1	1002	1207
57	234137.10.dec 3746685H1	47	317	57	234137.10.dec g3336428	1187	1565
57	234137.10.dec 6265607H1	93	216	57	234137.10.dec g3231388		
57	234137.10.dec 2503636H1	97	332			1189	1560
57				57 57	234137.10.dec 1575462T6	1191	1520
	234137.10.dec 698952H1	114	199	57	234137.10.dec 2585733H1	643	887
57	234137.10.dec.699726H1	114	366	57	234137.10.dec 1440773H1	645	908
57	234137.10.dec 699737H1	114	362	57	234137.10.dec g4075131	1202	1560
57	234137.10.dec 3051736H1	1008	1301	57	234137.10.dec 3503321H1	1200	1506
57	234137.10.dec 3789688H1	1025	1238	57	234137.10.dec g678032	1248	1560
57	234137.10.dec 389357H1	1031	1178	57	234137.10.dec 454365H1	1248	1428
57	234137.10.dec 2653885T6	1030	1522	57	234137.10.dec 3121619H1	1253	1545
57	234137.10.dec 389357R1	1034	1455	57	234137.10.dec 2867481H1	548	852
57	234137.10.dec 389357R6	1034	1426	57	234137.10.dec 3804490H1	551	852
57	234137.10.dec 4000387H1	1047	1317	57	234137.10.dec 5269079H2	559	779
57	234137.10.dec 1446567T6	1049	1521	57	234137.10.dec 5586156H1	563	775 725
57	234137.10.dec 4000387R6	1047					
57			1422	57 57	234137.10.dec 4711212H1	564	817
	234137.10.dec 4000387T6	1048	1526	57	234137.10.dec 3165089H1	566	832
57	234137.10.dec g2932747	1119	1560	57	234137.10.dec 3925365H1	677 .	960
57	234137.10.dec g4149909	1129	1566	57	234137.10.dec 5900267H1	666	950
57	234137.10.dec g3678702	1133	1563	57	234137.10.dec 5066375H1	690	941
57	234137.10.dec g2322637	1141	1560	57	234137.10.dec 5204570H1	708	961
57	234137.10.dec 644000H1	595	862	57	234137.10.dec 6111563H1	708	1024
57	234137.10.dec 4637553H1	573	818	57	234137.10.dec 3867362H1	706	960
57	234137.10.dec 3766620H1	627	862	57	234137.10.dec 1440773F6	645	1003
57	234137.10.dec 1742623H1	1002	1271	57	234137.10.dec 2736033H1	645	887
57	234137.10.dec 1742603H1	1002	1278	57	234137.10.dec 2948070H1	659	927
57	234137.10.dec 1742570H1	1002	1286	57	234137.10.dec 3872678H1	664	
57	234137.10.dec g1004759	1003	1262	57			952
57		628			234137.10.dec 1482169H1	22	287
57	234137.10.dec 3124271H1		935	57 57	234137.10.dec 1602692H1	23	223
	234137.10.dec 993448H1	632	867	57	234137.10.dec g775373	24	309
57	234137.10.dec 3875672H1	639	928	57	234137.10.dec 3530615H1	1	283
57	234137.10.dec g1813123	1485	1563	57	234137.10.dec 4171108H1	1	279
57	234137.10.dec g2631629	1507	1560	57	234137.10.dec 3156905H1	1	163
57	234137.10.dec 4077622H1	18	292	57	234137.10.dec g3191632	808	1227
57	234137.10.dec 3672287H1	17	227	57	234137.10.dec 2129471H1	807	1082
57	234137.10.dec 905762H1	20	222	57	234137,10.dec 3517838H1	819	1133
57	234137.10.dec g953765	20	379	57	234137.10.dec 5081676H1	841	1007
57	234137.10.dec 2816025H1	21	319	57	234137.10.dec 3870236H1		
57						846 1275	1114
	234137.10.dec g1815065	. 4	440	57 57	234137.10.dec g3896087	1375	1560
57 57	234137.10.dec g884950	8 .	411	57	234137.10.dec g5638159	1374	1566
57	234137.10.dec 3331405H1	13	269	57	234137.10.dec 589607H1	958	1206
57	234137.10.dec 6408514H1	16	601	57	234137.10.dec 1446567H1	968	1143
57	234137.10.dec 901600H1	430	735	57	234137.10.dec 1446567F6	968	1362
57	234137.10.dec 901500R1	430	955	57	234137.10.dec g1496263	983	1304
57	234137.10.dec 901744H1	430	628	57	234137.10.dec 1832689T6	986	1522
57	234137.10.dec 5591962H1	449	578	57	234137.10.dec 2287212H1	999	1230
57	234137.10.dec 2653885H1	511	789	57	234137.10.dec 5190833H2	125	366
57	234137.10.dec g868790	1436	1571	57			
		1730	13/1	3/	234137.10.dec 2723787H1	123	373

		Tal	ble 2 co	nt	,	PCT/US	800/25610
57 234137.10.dec 4880881	H1 127						
234137.10.dec 6245793	H1 105			7 234137.1	0.dec g269540	1 12	24 4500
234137.10.dec 2419654	H1 150			. 20413/.]	U.DEC ARRADAE	7 40	34 1560 34 1566
234137.10.dec 2121623	E6 110		5	23413/.7	0.dec a38077a	6 40	
27 234137.10.dec a47630a	8 440		_	234137.11	0.dec 2653885	6 12	44 1566
27 434137.10.dec n287576	8 110		5	7 234137.10	0.dec 4950523	F6 51	
57 234137.10.dec g498580	6 111		5	7 234137 10	0.dec 5091884	H1 51	
57 234137.10.dec 3327435	3 112	0 1566	57	7 234137 10	.dec 639236H		
57 234137 10 doc 1930000	H1 208	461	57		7.dec 639236H	1 52	777
57 234137.10.dec 1832689 57 234137.10.dec 1832689	H1 235	502	57		dec 1964688	H1 544	
- TOTAL OLUCE TRANSPORT	R6 235	564	57		.dec 53288851	11 783	
''O'. 'O.UEC /11591L	11 239	438	57 57		.dec 5782346F	11 700	
''O'. 10.UEC .331 /RAEL	11 300	546		234137.10	.dec 842587p1	907	
234137.10.dec a1616066	200	598	57	234137.10	.dec 8425g7⊔1	807	
234137.10.dec 2185641L	11 . 04-	605	58	480630.4.c	ec 1696097L	11 613	
204 137.10.dec 1618067L	11 00-		58	480630.4.d	ec 4922014H	1 628	- · ·
23413/.10.dec 02141025	23	599	58	480630.4.d	ec 978050H4		887
234137.10.dec d616022	25	441	58	480630.4.d	ec 3770059H	639	983
27 234137.10.dec a705506		291	58	480630.4.d	ec 4701108H	1 646	965
23413/.10.dec n705505	26	316	58	480630.4.d	ec 5278092H		923
57 234137.10.dec 4159960H	26	285	58	480630.4.de			855
57 234137.10.dec g831177		277	58	480630.4.de		663	886
57 234137.10.dec g574383	30	382	58	480630.4.de		666	733
57 234137.10.dec g868789	29	350	58	480630.4.de		667	915
	30.	312	58	480630.4.06		968	1178
	30	330	58	480630.4.de		997	1270
	32	435	- 58	480630.4.de		1172	
	23	311	58	480630.4.de		1187	1400
' ' O' . 10.UBC / AMAGONU1	34	305		480630.4.de	C 1334076LI1	1196	
204137.10.dec 288850784	34	96	58	480630.4.de	C 239876∩ы₁	1059	1425
204137.10.dec g814646	34	314	58	480630.4.de	C 3696304LI1	1065	1307
234137.10.dec 3243071H1	34		58	480630.4.de	01991868		1334
234137.10.dec 3511080H1	38	273	58	480630.4.de	20103211	1089	1473
234 137.10.dec 4518458H1		329	58	480630.4.ded	g1874953	1093	1497
23413/.10.dec 6496453U1	872	1113	58	480630.4.ded	201032R1	1094	1490
57 234137.10.dec 4344079H1	885	1320	58	480630.4.dec		1094	1632
57 234137.10.dec 4230253H1	906	1194	58	480630.4.ded		1094	1460
57 234137.10.dec g4152672	911	1192	58	480630.4.dec		1094	1329
57 234137.10.dec 5518712H1	953	1290	58	480630.4.dec		1100	1293
57 234137.10.dec 1255603H1	409	640	58	480630.4.dec		1100	1664
57 234137 10 dec 1255603H1	410	653	58	480630.4.dec		1101	1337
	410	762	58	480630.4.dec		1100	1333
		720	58	480630.4.dec		1117	1261
- TOULUEC ONATURE	4.0.0.	1560	58	480630.4.dec	6074887H1	1118	1418
1.07.10.dec 0.17019gg		1569	58	480630.4.dec	5442484H1	1120	1342
234137.10.dec g2107226		1560	_	480630.4.dec	435271H1	1130	1338
234137.10.dec 2691496T6		1534	58 50	480630.4.dec	1400593H1	1145	1339
27 234137.10.dec g8145g6	444-	575	58	480630.4.dec	1800427H1	1160	
37 234137.10.dec g2782755	4	508	58	480630.4.dec	601700H1	1199	1430
23413/.10.dec 04327424			58	480630.4.dec	2022558H1	1215	1448
234137.10.dec 01516847		575 570	58	480630.4.dec	5223360H1		1474
234137.10.dec g884865		579	58	480630.4.dec	5098384H1	1218	1485
234137.10.dec g831967		574 570	58	480630.4.dec	2306363	1276	1500
37 234137.10.dec g287480g		576	30	480630.4.dec	6078781H1		1541
57 234137.10.dec 01406164		525	58 .				1606
37 434137.10.dec d388300E		560	58	480630.4.dec	92343652		1728
57 234137.10.dec 4772565H1		569		480630.4.dec	5515833H1	1320	1615
57 234137.10.dec g822266	1258 1	527			5604966H1	1328	1577
	1282 1	569		180630.4.086	4535083T1		1904
''''''''''''''''''''''''''''''''''''		60		180630.4.dec	2562006H1		1621
	465.	60	- OO 4	100030.4.dec	2411212H1	4	559
- · · · · · · · · · · · · · · · · · · ·	4	63		80630.4.dec	1981090H1	4	611
· · · · · · · · · · · · · · · · · ·	4000	66	58 4	80630.4.dec	1981090R6	4	858
234137.10.dec 04330570		66	58 4	00030.4.dec	1506522H1	40	
3/ 23413/.10.dec 04533800			58 4	80630.4.dec	1504651H1		567 614
3/ 23413/.10.dec a56664a		60 60	58 4	80630.4.dec 1	1981090T6	4004	614
234137.10.dec 02752755	1347 15		58 4	80630.4.dec 2	7446	4 44 -	893
3/ 23413/.10.dec a342422a	1348 15		58 4		0000000		655
3/ 23413/.10.dec 1964266µ1	1373 15		58 40	80630.4.dec 4	4 45756111		892
9/ 63413/ 10 dec 1061000==	1221 14			30630.4.dec 3	407000144		748
2/ 43413/ 10 dec 1064000===	1221 153	35	58 48	30630.4.080 3	407885H1		82
57 234137 10 dec e645444	1222 152			30630.4.dec 5 30630.4.dec g	443335H1		573
	1227 156				2141427 ·		88
	-			0630.4.dec 5	40000011.		15
		106				10	

	-		Table	2 cont.		•	
480630.4.dec	2793926H1	1433	1744	58	480630.4.dec	q1976684	808
480630.4.dec	g2343629	1432	1796	58	480630.4.dec		831
480630 4 dec	5490024H1	1435	1723	58	480630 4 dec	3177860H1	831

58 1241 58 1133 58 1152 58 480630.4.dec 2626277H1 1447 58 1673 480630.4.dec 925425H1 897 1202 58 480630.4.dec 3219835H1 1451 1757 58 480630.4.dec 924684R1 897 1331 g1390343 58 480630.4.dec 1450 1891 58 480630.4.dec 924684H1 898 1169 58 58 480630.4.dec 1803241H1 1454 1674 480630.4.dec 3179844H1 914 1225 480630.4.dec 58 480630.4.dec 4773772H1 1462 1725 58 5433521H1 949 1186 58 480630.4.dec 5174892H1 1468 58 480630.4.dec 1555 2553966H1 949 1223 58 1472 58 480630.4.dec 4367404H1 1725 480630.4.dec g1445465 967 1341 58 . 1475 480630.4.dec g2837598 58 480630.4.dec 1932 5044309H1 423 693 58 58 480630.4.dec g4736639 1487 1938 480630.4.dec 424 676 2151905H1 58 2131706H1 480630.4.dec g2728681 1490 1932 58 480630.4.dec 450 720 58 g3756314 480630.4.dec 1511 1932 58 480630.4.dec 2868958H1 458 681 58 480630.4.dec q3330906 1508 1933 58 480630.4.dec 2106234H1 326 573 58 480630.4.dec 2758440H1 1514 1777 58 480630.4.dec 3296308H1 415 672 58 480630.4.dec 3296308T6 1517 58 1898 480630.4.dec 3296308F6 415 848 58 480630.4.dec g2657800 1516 1932 58 480630.4.dec g1645413 96 1 58 480630.4.dec g4568716 1527 1932 58 480630.4.dec 1732045H1 284 g3253868 58 480630.4.dec 1521 1933 58 480630.4.dec 87 331 2451711H1 58 480630.4.dec 1209430T1 1534 58 1896 480630.4.dec 6603078H1 170 599 58 480630.4.dec g4618917 1534 1932 59 480951.5.dec 2613688H1 249 58 480630.4.dec g3445980 1534 59 1932 480951.5.dec 587 1371886H1 367 58 480630.4.dec 201032F1 1550 1932 59 480951.5.dec 2741956H1 367 615 58 480630.4.dec 1834212H1 1559 59 1834 480951.5.dec 3574186H1 367 637 58 480630.4.dec 1834212T6 1559 1893 59 480951.5.dec 3745122H1 378 679 58 480630.4.dec g2219997 1572 1939 59 480951.5.dec 491592H1 380 531 815632H1 58 480630.4.dec 1579 1813 59 480951.5.dec 2441829H1 375 581 58 480630.4.dec g2538714 1581 1944 59 480951.5.dec 6100267H1 376 663 58 480630.4.dec g1388684 1586 1932 59 480951.5.dec 378 725 g1974420 g2195437 58 480630.4.dec 1593 1932 59 480951.5.dec 379 663 g1718741 58 480630.4.dec 2825773T6 1598 1890 59 480951.5.dec 2599377H1 366 634 58 480630.4.dec 1611 g2161259 1937 59 480951.5.dec 3331491H1 382 620 58 480630.4.dec 2460384H1 1611 1835 59 480951.5.dec 3336103H1 1167 1426 g4740936 58 480630.4.dec 1609 1940 59 480951.5.dec 3359017H1 1168 1449 58 g3213847 480630.4.dec 1615 1942 59 480951.5.dec 2508116H1 1180 1417 58 480630.4.dec q4892585 1622 1935 59 480951.5.dec 1420 5217308H1 1180 58 480630.4.dec g2115665 1637 1932 59 480951.5.dec 5733042H1 1217 1442 58 g315021 480630.4.dec 1659 1932 59 480951.5.dec 4115350H1 1217 1471 58 480630.4.dec g3422567 1665 1932 59 480951.5.dec 6315280H1 1435 1265 58 480630.4.dec 1665 g1241338 1932 59 480951.5.dec g2266304 1392 1539 58 480630.4.dec g2779212 1668 1898 59 680479H1 480951.5.dec 1445 1607 58 480630.4.dec g825084 1696 1942 59 480951.5.dec 3726996H1 403 691 58 480630.4.dec 1704 1943 59 g3277734 480951.5.dec 1989538H1 367 687 58 480630.4.dec 3706040H1 504 764 59 480951.5.dec 4374001H1 367 657 58 480630.4.dec 2726096H1 504 728 59 480951.5.dec 3385558H1 367 632 480630.4.dec 58 6606548H1 535 1097 59 480951.5.dec 3541978H1 18 225 58 5954134H1 480630.4.dec 4947185H1 569 799 59 480951.5.dec 806 1123 58 480630.4.dec 5202184H1 607 806 59 480951.5.dec 2780282H1 808 970 g3889620 58 1746 480630.4.dec 5212745H1 1939 59 480951.5.dec 848 1017 58 480630.4.dec g1875009 1758 1940 59 480951.5.dec 6570768H1 868 1329 58 480630.4.dec g2732264 1818 1942 59 480951.5.dec 2252058H1 1002 1224 g2007592 59 58 480630.4.dec 1851 2165 1498 480951.5.dec 2020306T6 995 g2898362 58 480630.4.dec 1863 1932 59 480951.5.dec 6110583H1 1002 1204 g3047837 58 480630.4.dec 1869 1932 59 480951.5.dec 2721730H1 1002 1213 58 480630.4.dec 4947804F6 773 1232 59 480951.5.dec 338780H1 1002 1193 58 480630.4.dec 1693713H1 1002 773 59 480951.5.dec 3527343H1 1008 1285 58 480630.4.dec 3576689H1 780 1074 59 480951.5.dec 6568044H1 1080 1385 58 480630.4.dec 548416H1 59 781 921 480951.5.dec 1098 2128264H1 1377 58 480630.4.dec 4992743H1 747 944 59 480951.5.dec 1103 1361 679425H1 58 480630.4.dec 4947804H1 773 874 59 480951.5.dec 1108 1351 630229H1 58 480630.4.dec q1986962 673 940 59 480951.5.d c 2126237H1 1115 1363 480630.4.dec 4535083H1 58 670 804 59 480951.5.d c 1115 1380 4060761H1 58 480630.4.dec g4929642 674 1944 59 480951.5.dec 546732H1 1115 1359 58 480630.4.dec g792010 696 1021 59 480951.5.d c 378369H1 1122 1360 4992948H1 58 480630.4.dec 747 1021 59 480951.5.dec 451941H1 1124 1325 58 480630.4.dec 4992572H1 747 1009 59 480951.5.d c 4380896H1 1324 1126 5557064H1 58 480630.4.dec 747 59 1005 480951.5.dec 510978H1 1144 1360

	59 48095	4 5 4			Tab	le 2 con	t.) 	PCT/U	500/25610
		1.5.dec 1 1.5.dec 1	648031H1	1149	1349	59					
	.0000		648008H1	1149	1360	59		1.5.dec		H1 36	896
	59 48095	1.5.dec 4	044321H1	1150	1385	59		.5.dec	1346252	H1 37	
5	9 48095		216861H1	1150	1417	59		.5.dec	3470714	H1 54	
5	9 480951		30261H1	1165	1471	59	,	.5.dec		H1 55	
5	9 480951		2595H1	1166	1383	59	.00001	.o.dec		1 55	5 733
5	9 480951		79085H1	394	573	59	480951	.5.dec		H1 55	9 837
5	9 480951		42261H1	394	657	59		.5.dec		H1 55	8 810
5	y 480951	.5 dec 32	147442 47844H1	397	721	59	480951	.5.dec	94310946	60	
5	9 480951.		47644H1 41674H1	409	700	59	480951.	5.dec	3111016	11 608	
59	9 480951.	.5.dec 31.	40608H1	408	713 .	. 59	480951.	5.dec	g3178543	603	960
59	480951.	5.dec a16	972153	409	691	59	480951.	5 dec	452965H	604	819
59	480951.	5.dec 613	367340	410	646	59	480951.	5 dec	g2056998		223
59	48 0951.	5.dec 55/	15462H1	408	805	59	480951.	5 dec	6322880H		
59	480951.	5.dec - 014	25731		605	59	480951.5		999590H1 3758338H		
59	480951.	5.dec 109	8757R6		900 :	59	480951.5		3326928H		
59	480951.	5.dec 198	8757H1		796	59	480951.5		2729711H		670
59	480951.5	5.dec - 010	78134		686	59	480951.5	.dec	g1146694		658
59 50		5.dec 149	^^-		714	59	480951.5	.dec	2020306F6	403	748
59 59		o.dec g10	68989		541 740	59	480951.5	.dec :	2133705H	403	796
59	480951.5	o.dec 266	4883H1	•	551	. 59	480951.5	.dec :	3647806H1		657
59 59	480951.5	.dec 253	5301H1	-	346	59	480951.5	dec 4	4822653H1		718
59	480951.5	.dec g131	13751		20	59	480951.5	dec d	1625996	401 403	664
59	480951.5	.dec 1236	075F1	'	050	59 50	480951.5.	dec 4	842127H1	402	763
59	480951.5		049H1 :		55 55	59	480951.5	dec 2	2020306H1	403	660
59	480951.5. 480951.5.	.dec 3571	943H1 4		75	59 59	480951.5.	dec 4	418643H1	401	681 662
59	480951.5.		291H1		73	59 59	480951.5.	dec 4	747674H1	401	673
59	480951.5.		<u>49</u> H1 3		31	5 9	480951.5.0	1 8 C 4	174839H1	401	682
59	480951.5.		779H2 3		32	59	480951.5.0	sec 5	395040H1	393	657
59	480951.5.			70 63		59	480951.5.0	iec 5	542605H1	392	605
59	480951.5.0			71 57		59	480951.5.0	iec 3	117868H1	277	467
59	480951.5.0		· · · ·	72 63	13	59	480951.5.d		243270H1	307	551
59	480951.5.0			52 59	5	59	480951.5.d 480951.5.d		750596H1	1	162
59	480951.5.0	dec 39873 dec 42051		52 62	9		480951.5.d		78980H1	1	467
59	480951.5.d	ec 25204					480951.5.d	ec gz	229512	237	648
59	480951.5.d	ec 28786	40H1 3€			59	480951.5.de	ec 55	42378H1	251	437
59	480951.5 d	ec 41701				59	480951.5.de	sc 33	95687H1	241	487
59	480951:5.d	ec 36400				59	480951.5.de	oc 65	95818H1 62968H1	242	533
59	480951.5.de	BC 33631	34H1 37 30H1 37			59 .	480951.5.de		02908H1 15741H1	290	812
59 50	480951.5.de	ec 20476	16H1 37			59 4	480951.5.de	C 631	80347H1	271	525
59 59	480951.5.de	9 C 687508	3H1 14.			59 4	480951.5.de	c 37	51944H1	369	685
59 59	480951.5.de	€ 223514	8H1 33			39 4	180951.5.de	C 478	36273H2	368	569
59	480951.5.de	€ 489142	4H1 336			59 4	180951.5.de	0 350)4819H1	368	548
59	480951.5.de	⊬ g20130	66 33/			59 4	180951.5.de	011	98636	369 372	671
	480951.5.de		8F6 335			59 4	80951.5 da	g20	25536	372	697 650
	480951.5.de 480951.5.de		11 335	515		59 4 59 4	80951.5.dec		1711H1		658 670
	480951.5.de		3H1 336	922			80951.5.ded	357	9019H1		670
	480951.5.dec		DH1 340	632		•	80951.5.ded	g89:	2899		988 929
	480951.5.dec			551			80951.5.dec		7937H1	`	926
59 4	480951.5.dec			701		59 48	80951.5.dec	3762	204H1		967
59 4	480951.5.dec	8550201		1198	1		30951.5.dec 30951.5.dec		1346H1	686	943
-	100331.5.dec	5200000		853			30951.5.dec		2679H1		31
59 4	180951.5.dec	0/10/00	_	874			10951.5.dec		053H1	-	72
59 4	180951.5.dec	0802010		935			0951.5.dec		432H1	722 9	70
59 4	80951.5.dec	1660116		1006			0951.5.dec			732 9	87
59 4	80951.5.dec	9867950		855	:	59 48	0951.5.dec			733 9	86
59 4	80951.5.dec	9867850		836			0951.5.dec	3291		743 9	70
59 4	80951.5.dec	g1687504		1061			0951.5.dec	g1303		752 11	022
59 4	80951.5.dec	1660036		1039			0951.5.dec	39847		751 92	28
59 40	80951.5.dec	4459507F		763	5	9 480	0951.5.dec	36788		7 59 97	
59 48	30951.5.dec	2243455h		888			951.5.dec	34416		60 97	
59 4 8	30951.5.dec	2737280H		901	5	9 480		36748	FOLIA	61 94	
59 48	30951.5 dec	3051410H		876	5	9 480	951.5.dec	19090	02H] 7	62 97	
59 48	0951.5.dec	650429H1	380	946	5	9 480	951.5.dec	33221 75403			64
59 48		5224648H	380 1 381	614	5	9 480	951.5.dec	75403 17271		93 97	0
		• . •	. 301	<u>.</u> 481	- 59	9 480		55160		93 97	
					108			-0.00	1001 7	93 97	0



Table	2	cont.
614		60

				1 abi	e z com.				
59	480951.5.dec	5589241H1	390	611	60	350399.5.dec	5119208H1	31	322
59	480951.5.dec	g2166364	390	767	60	350399.5.dec	4968666H1	38	167
59	480951.5.d c	1236401F6	443	871	60	350399.5.dec			
59	480951.5.dec	5577131H1	448	701	60		4956016H1	18	120
59			_			350399.5.dec	4515864H1	21	216
	480951.5.dec	g1163670	449	608	60	350399.5.dec	3215683H1	25	294
59	480951.5.dec	2779615H1	475	732	60	350399.5.dec	4511482H1	1566	1822
59	480951.5.dec	3211910H1	477	597	60	350399.5.dec	2909383H1	1567	1828
59	480951.5.dec	5615524H1	501	808	60	350399.5.dec	g4392449	1572	1941
59	480951.5.dec	2705619H1	492	753	60	350399.5.dec	6023589H1	1586	1855
59	480951.5.dec	2544567H2	526	783	· 60	350399.5.dec		1627	1852
59	480951.5.dec	1592175H1	534	730	60	350399.5.dec	4824771H1	1318	1566
59	480951.5.dec	4176769H1	535	710	60	350399.5.dec	g3678597	1337	
59	480951.5.dec	1591703H1	534	728	60		•		1727
59	480951.5.dec					350399.5.dec	2486926H2	1386	1624
59		g1278591	492	637	60	350399.5.dec	1289520T6	3923	4170
	480951.5.dec	g1156402	537	763	60	350399.5.dec	1289520F6	3923	4219
59	480951.5.dec	g1670312	491	742	60	350399.5.dec	94109131	3931	4211
59	480951.5.dec	g1494000	492	657	60	350399.5.dec	4351280H1	3943	4211
59	480951.5.dec	g1721860	492	889	60	350399.5.dec	4432226H1	4 -	262
59	480951.5.dec	g989068	501	757	60	350399.5.dec	6380386H1	8	319
59	480951.5.dec	3800372H1	367	651	60	350399.5.dec	5498050H1	15	259
59 .	480951.5.dec	2448255H1	367	598	60	350399.5.dec	1860316H1	1270	1490
59	480951.5.dec	3765155H1	367	676	60	350399.5.dec	g1062524	3959	4199
59	480951.5.dec	6374408H1	367	606	60	350399.5.dec	_		
59	480951.5.dec	5538463H2	366	555				3971	4216
59	480951.5.dec				60	350399.5.dec		3972	4213
		2921720H1	380	641	60	350399.5.dec	g4888572	3975	4211
59 50	480951.5.dec	g2057109	160	442	60	350399.5.dec	g3155094	3986	4213
59	480951.5.dec	6481690H1	361	892	60	350399.5.dec	g1062503	3996	4186
59	480951.5.dec	595420H1	362	604	60	350399.5.dec	1602855H1	184	379
59	480951.5.dec	3591022H1	361	660	60	350399.5.dec	2716719H1	184	433
59	480951.5.dec	1320037H1	362	625	60	350399.5.dec	1300839T6	1821	2141
59	480951.5.dec	3593512H1	362	680	60	350399.5.dec	g5366860	1838	2179
59	480951.5.dec	5391147H1	362	657	60	350399.5.dec	g5630541	1865	2172
59	480951.5.dec	2719839H1	362	612	60	350399.5.dec		1537	1846
59	480951.5.dec	g1277584	354	847	60	350399.5.dec	355153H1	1553	1718
59	480951.5.dec	2457980H1	359	591	60	350399.5.dec	g4833913	3956	4213
59	480951.5.dec	g1799076	359	819	60				
59	480951.5.dec	5538138H1				350399.5.dec	4750324H1	972	1083
5 9			361	516	60	350399.5.dec	g872980	998	1323
	480951.5.dec	4023870H1	342	601	60	350399.5.dec	1300839H1	897	1172
59	480951.5.dec	1360005H1	342	581	60	350399.5.dec	5515034H1	915	1111
59	480951.5.dec	g1998895	343	503	60	350399.5.dec	097987H1	926	975
59	480951.5.dec	4161775H1	343	585	60	350399.5.dec	g839057	935	1323
59	480951.5.dec	3778265H1	342	647	60	350399.5.dec	2553277H1	796	1058
59	480951.5.dec	4843784H1	345	615	60	350399.5.dec	g2838628	810	1289
59	480951.5.dec	3533957H1	347	657	60	350399.5.dec	3489831H1	859	1131
59	480951.5.dec	4520442H1	349	604	60	350399.5.dec	g3148335	879	1034
59	480951.5.dec	g2669182	350	780	60		1300839F6	897	1342
59	480951.5.dec	4618672H1	349	592	60	350399.5.dec	g4486333	899	1289
59	480951.5.dec	4031302H1	342	592	60	350399.5.dec			
59	480951.5.dec	3321371H1	354	632				4053	4211
59	480951.5.dec	4729423H1			60	350399.5.dec	3560750H1	4026	4206
			355	513	60	350399.5.dec	3	4054	4212
59	480951.5.dec	2581094H1	355	588	60	350399.5.dec		4079	4211
59	480951.5.dec	3250802H1	341	642	60	350399.5.dec	g1812272	4113	4376
59	480951.5.dec	2913426H1	342	600	60	350399.5.dec	3899577H1	1	168
59	480951.5.dec	1360005F1	342	828	60	350399.5.dec	2546733H1	1	228
59	480951.5.dec	4770833H1	341	602	60	350399.5.dec	3205796H1	1 .	128
59	480951.5.dec	5395039H1	366	630	60	350399.5.dec		1	222
59	480951.5.dec	4067217H1	365	632	60	350399.5.dec		333	619
59	480951.5.dec	4160969H1	364	644	60	350399.5.dec		338	623
59	480951.5.dec	3538760H1	363	573	60	350399.5.dec			
59	480951.5.dec	5064082H1	364	575	60			1126	1356
59	480951.5.dec	407591H1				350399.5.dec		1135	1392
			363	520	60	350399.5.dec	0	1140	1306
59	480951.5.dec	5542680H1	363	579	60	350399.5.dec	g692250	1149	1297
5 9	480951.5.dec	3394195H1	364	627	60	350399.5.dec	1579183H1	1248	1423
59	480951.5.dec	2885143H1	. 364	622	60	350399.5.dec	1274218H1	1425	1658
60	350399.5.dec	3889877H1	117	388	60	350399.5.dec	g1062502	1438	1700
60	350399.5.dec	6219217H1	155	255	60	350399.5.dec	g1062523	1439	1733
60	350399.5.dec	g1784403	102	397	60	350399.5.dec	5510701H1	1451	1542
60	350399.5.dec	g1665816	26	4211	60	350399.5.dec	5407906H1	1875	2083
		-	-		100				

		Table 2		PC	T/US00/25610
60 350399.5.dec 279320	72114 4000	Table 2	ont.		
60 350399.5.dec 500686		00	60 350399.5.de	A63004444	
60 350399.5.dec g31907		2040	60 350399.5.de		3395 3650
60 350399.5.dec 186538		2190	60 350399.5.de		3418 3605
60 350399.5.dec g28339		2224	60 350399.5.de		3468 3754
		2179	60 350399.5.de		3473 3694
		2135	60 350399.5.de		3473 3862
		2183	60 350399.5.de		3473 3747
		2183	60 350399.5.de		3475 3781
		2421	60 350399.5.dec		3483 3930
	3H1 2128	2403	60 350399.5.dec		3505 3789
	H1 2956	3201	60 350399.5.ded		3516 3787
	¹⁴ 2219	0.400			3520 3757
	H1 2219				3524 3748
	2276	0.455			3541 3787
	7 2312	070 /		1266266H1	3541 3774
60 9181259	6 2325	0400			3575 3919
50 050000.0.000 2083116	H1 2324	0010			3577 4025
60 900000.0.000 91812/5	6 2326	0505			3580 3848
50 Journal 2008841	H1 2326	0570		2447927T6	3593 4173
	H1 2333	0004		2292715T6	3592 4172
50 January 1719208	-6 2402	0746		5059812H1	3597 3907
	11 2412	0700		6394265H1	3604 3820
60 2824/32	11 2412	2712 6		2814355T6	3639 4168
2289458	11 240e -	2627 6		5877319H1	3644 3917
207/589	11 2508 4	2788 6		4087702H1	3647 3928
CO 000000.0.UEC 1/19208F	11 2510 4	2712 6	,	1005310H1	3665 3934
60 0000000 453/301H	1 2540	2793 66	J.uec	5861292H1	3741 4031
3290129H	1 2542	2783 60	000000.0.066	1859304T6	3749 4174
	2558 -	2782 60	444000.0.060	94650781	3779 4209
20 20000.0.080 95363819	2561 3	712 60	355555.5.060	g3742022 :	3779 4220
	1 2508 2	1152 60	22200.0.000	g4264942 :	3790 4211
60 05005.5.dec 4920360H	1 2650 2	930 60		5206350H1 3	798 4021
	1 2702 2	902 60	00000.0.066	g4738038 · 3	799 4209
CO	1 2750 o	824 60	350399.5.dec	g3649493 g	803 4220
	2765 2	328 60	350399.5.dec	1431740T6 3	808 4168
	2765 2	073 60	350399.5.dec	93539316 a	812 4215
	2970 2	238 60	350399.5.dec	3539304 3	812 4213
CO 200000.0.UBC 20434/3H1	2795 20	063 60	350399.5.dec	1637463T6 3	822 4170
20 0411224H1	3034 25	307 60	350399.5.dec 5	5949046H1 3	823 4074
		950 60	350399.5.dec 1	620983T6 3	330 4190
	2803 34	06 60	350399.5.dec 1	637463F6 3/	329 4183
1431/4086		37 60	350399.5.dec 1	637463H1 3	329 4043
CO 1431/40H1	2862 30		350399.5.dec g	1549797 38	330 4211
60 350399.5.dec 1431740R1 60 350399.5.dec 2447927H1	2862 31		350399.5.dec g	2069536 <u>3</u> 8	29 4209
CO 2000.0.0EC 244/92/H1	2942 31		350399.5.dec g	2942270 - 38	29 4216
	2942 34		350399.5.dec g	4186986 38	30 4211
CO 3446015H1	2954 30		350399.5.dec g	2942265 3a	30 4211
20 20000.5.dec 53/2158H1	2955 318		350399.5.dec g	3250503 38	
60 1019598H1	2956 315		350399.5.dec g	1114971 38	
60 350399.5.dec 664175H1	3037 328		350399.5.dec g3	3092120 38	
60 0408282H1	3043 356		350399.5.dec g4	1089641 3a	
60 -00005.5.060 912/1956	3060 341		350399.5.dec g4	175757 383	
CO 3134408H1	3061 333		350399.5.dec 94	08/874	
60 00000.0.dec 3255616H1	3104 336		350399.5.dec 94	0848R1 384	
60 250053.5.06C 4089831H1	3182 345	_	350399.5.dec 94	0848H1 386	
CO 93255048	3211 359	_	350399.5.dec g9	07438 386	
60 05000.0.06C 0352165H1	3213 331	_	350399.5.dec 17	40387T6 385	
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350399.5.dec 4934215H1	3219 3460		330399.5.dec na	558706 387	
350399.5.dec 2459559H1	3247 3476		350399.5.dec 959	9456R1 388	
50 350399.5.dec 1620983F6	3265 3745		350399.5.dec 950	456H1 388	
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350399.5.dec 5103592H1			350399.5.d c 642	0040-	
550399.5.dec 1859304F6			350399.5.dec	7	
350399.5.dec 1859303H1			350399.5.d c 128		
350399.5.dec 6614830H1			350399.5.dec 304		
350399.5.dec 1894769H1			350399.5.dec 147		
60 350399.5.d c 4630494H1		60	350399.5.dec 147	4400114	
	3395 3655	60.			
		110		663T6 1496	1669
		-			

Table 2 cont. 350399.5.dec 3941003H1 085713.2.dec 3927329H1 350399.5.dec 4672530H1 085713.2.dec 2593752H1 350399.5.dec g1471517 085713.2.dec g1981200 350399.5.dec g751699 085713.2.dec 2097957H1 350399.5.dec g692289 085713.2.dec 3769037H1 350399.5.dec g714847 085713.2.dec 1683135H1 350399.5.dec 112663R6 085713.2.dec 3509114H1 350399.5.dec 112663H1 085713.2.dec 5832853H1 350399.5.dec 3719539H1 085713.2.dec 806178H1 2844152H1 350399.5.dec 085713.2.dec 3558496H1 350399.5.dec 5347842H1 085713.2.dec 2795155H1 350399.5.dec 6494421H1 085713.2.dec 1982603R6 q3003587 350399.5.dec 085713.2.dec 2902677H1 350399.5.dec 1274218F1 .1425 085713.2.dec q4451382 350399.5.dec 3409696H1 085713.2.dec g316154 350399.5.dec g3645371 085713.2.dec 5819630H1 350399.5.dec 4532952H1 085713.2.dec 2869957H1 350399.5.dec 1879147H1 085713.2.dec 2869957F6 350399.5.dec 3719622H1 085713.2.dec g942739 350399.5.dec g3425750 085713.2.dec g1477126 350399.5.dec q3213638 085713.2.dec g1527541 g1784345 350399.5.dec 085713.2.dec g969342 350399.5.dec 4376792H1 085713.2.dec g2184408 350399.5.dec 4376541H1 085713.2.dec 5307452H1 350399.5.dec 4128636H1 085713.2.dec 6321277H1 350399.5.dec g566326 085713.2.dec 2869957T6 350399.5.dec g751651 085713.2.dec g2037747 350399.5.dec g814715 085713.2.dec 2463542H1 350399.5.dec 5095975H2 085713.2.dec 2463542F6 g1471430 350399.5.dec 085713.2.dec 2044369H1 350399.5.dec 1382644H1 085713.2.dec 1772731H1 350399.5.dec 6387138H1 085713.2.dec 4266001H1 350399.5.dec 5833550H1 085713.2.dec 4730142H1 350399.5.dec 2723562H1 085713.2.dec g2555947 350399.5.dec 5413045H1 085713.2.dec 840781H1 350399.5.dec 3451343H1 085713.2.dec 4593969H1 085713.2.dec 2876519H1 085713.2.dec g5109339 085713.2.dec g3804453 085713.2.dec 5821366H1 085713.2.dec 4714518H1 085713.2.dec 5818829H1 085713.2.dec g4522739 085713.2.dec 5813545H1 085713.2.dec g5369593 085713.2.dec 5817650H1 085713.2.dec 5563230H1 085713.2.dec 5812917H1 085713.2.dec 5327933H1 085713.2.dec 6518401H1 085713.2.dec q3658839 085713.2.dec 2655535T6 085713.2.dec g3927414 085713.2.dec 5910362H1 085713.2.dec 5275308H1 085713.2.dec 2878073H1 085713.2.dec 2673389H1 085713.2.dec g944173 085713.2.dec 5333974H1 085713.2.dec 3702759H1 085713.2.dec 1982603H1 085713.2.dec g5325988 085713.2.dec 4726834H1 085713.2.dec g969343 085713.2.dec g4196655 085713.2.dec 2043940H1 085713.2.dec 3484912H1 085713.2.dec 661586H1 085713.2.dec 2552885T6 085713.2.dec 5056769H1 085713.2.dec 552697H1 085713.2.dec 6306802H1 085713.2.dec g2594419 085713.2.dec 3042074H1 085713.2.dec g1647965 085713.2.dec 4261452H1 085713.2.dec 613645H1 085713.2.dec 2773951H1 085713.2.dec 2132842H1 085713.2.dec g3050704 085713.2.dec g4073710 085713.2.dec 6023537H1 085713.2.dec g1527498 085713.2.dec 2653867H1 085713.2.dec 5412976H1 085713.2.dec g2184179 085713.2.dec 2721848H1 085713.2.dec g3960542 085713.2.dec 2637714H1 085713.2.d c g1119058 085713.2.dec g274328 085713.2.dec 6411539H1 3801333H1 085713.2.dec 085713.2.dec g3077101 085713.2.dec 2857263F6 085713.2.dec q2958693 085713.2.dec 2857263H1 085713.2.dec 3640567H1 085713.2.dec 6212439H1 085713.2.dec g2569732

						T- 1						PCT/US	500/2	5610
	61	085713.2	dec 225700	01.14		lab	le 2 co	ont.			,			2.010
	61	085713.2	.dec 226728 .dec 581463	4114		1550		63	117464.	7 dec	GE60004			
	61	085713.2.	d c 581764			1515		63	117464.	7.dec	g563294 6247611	9 20	266	2169
	61	085713.2.	dec 582246		4	1517	(63	117464.	7.dec	689848R	H1 1		333
	61	085713.2.	dec 581677			1441		63	117464.7	7.dec	6482192	6 1 41 10		234
	61	085713.2.	dec 042950			1504		63	11/404./	dec.	8517510			501
	61	085713.2.0	dec 2463542			1388		53	117464.7	dec.	54633491	1 21 11 39	_	828
	61	085713.2.0	dec 0198216			1724		53	117464.7	'.dec	9317507	,, 39 50	_	522
	61	085713.2.0	dec 0867230			526		33	117464.7	.dec	4677105H	11 67		855
	61	085713.2.0	dec 0147703	_ ~		2737 2738		3	117464.7	.dec	5541478H	11 72:		937 938
	61	085713.2.0	iec 646057h	11 2	_	739		3	117464.7	.dec	4028742H	11 869	_	1131
	61 61	085713.2.0	iec 6158466	H1 2		647		3	117464.7	.dec	3966996H	1 932		1103
	61	085713.2.d	lec 3535428	H1 25		737	6		117464.7.	.dec	3967229H	1 931		1163
	61	085713.2.d	lec g147935.	4 25		736	6: 6:		117464.7.	dec	3966996F	6 931		1419
	61	085713.2.d		3 25		661	63		117464.7.	dec	1843101H	1 966		244
•	61	085713.2.d 085713.2.d		1 1 37		76	63		117464.7.		1843101R	6 966		411
	61	085713.2.d		11 41		50	63		117464.7. 117464.7.		5989305H	1 101	_	302
	61	085713.2.de		48	5 75	54	63		117464.7.		3532784H	1 113		395
	61	085713.2.de		• •		54	₁ 63		117464.7.		3518507H1		2 1	539
	61	085713.2.de	ec 2655535H ec 2655535F				63		117464.7.0		3876438H1	. : :	•	591
	61	085713.2.de	2655535F	_			63	}	117464.7.0		1996445H1 722076H1		_	590
	61	085713.2.de	C 1902994H				63	}	117464.7.0		3162945H1	142	•	639
	62	245014.1.de	C 6345408H	1 336			63		117464.7.0		g1925687			799
	62	245014.1.de	C 4435621H	1 611 1 722			63		117464.7.d	iec 4	4517215H1	1530		954
	62	245014.1.de	C 4439530H	1 634		_	63		117464.7.d	lec d	9751107	1557 1572		311
	62	245014.1.de	C 4972420H	1 680	•		63	•	117464.7.d	ec :	5044574H1	1597		774
	62	245014.1.de	C 3275925H	1 732			63	1	117464.7.d	ec 3	3966996T6	1634	20	881 219
	62 62	245014.1.de	c g3446618	733			63	1	17464.7.d	ec 6	3514359H1	1648		85
	62 62	245014.1.dec		733			63]	17464.7.d	ec 8	351751T6	1661		∞ 79
	62	245014.1.dec		741	118		63 63	,	17464.7.de		411633H1	1707		30
	52 52	245014.1.dec 245014.1.dec		755	100		63	'	17464.7.de		556621H1	1743		87
		245014.1.dec		762	102		63	1	17464.7.de 17464.7.de		556364H1	1743	20	
		245014.1.dec			991		63	i	17464.7.de		005955H1	1755	19	
		245014.1.dec		871	126	1	63	1	17464.7.de		3253416	1795	22	17
6	2	245014.1.dec	2872604H1 4212104H1	874	113	5	63	1	17464.7.de		4300096	1792	220)4
	2	245014.1.dec	94901006	921	118		63	1	17464.7.de		049127F6 049127H1	1831	22€	
	2	245014.1.dec	960167141	896	956		63	11	17464.7.de		594085H1	1831	210	
6	2	245014.1.dec	3450361D6	1044 68		7	63	11	17464.7.de	c 58	370047H1	2627	288	
6	2 2	245014.1.dec	5399133H1	1	532		63	11	17464.7.de	c a2	2016814	2700 2540	294	
6:	2 2	245014.1.dec	3450361H1	68	210 323		63	- 11	7464.7.ded	c 85	1751H1	192	288	
62		245014.1.dec	2731452F6	81	415		63	11	7464.7.ded	68	9848T6	1650	443 212	
62 62		245014.1.dec	2731452H1	81	316		63	11	7464.7.ded	: 58	11873H1	2077	235	.
62	_	45014.1.dec	263691H1	94	421		63 63	11	7464.7.dec	: g7:	50993	1844	221	
62	_	45014.1.dec	5674201H1	106	342		63	11	7464.7.dec		9952H1	1871	211	
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Table 3 Tissue Distribution	Embayonic Starchuser, 179, Gorm Colle, 189, Cardiomedia 200,	Sense Organs - 44%, Unclassified/Mixed - 20%, Connective Tissue - 11%	Germ Cells - 29%, Digestive System - 12%, Embryonic Structures - 11%	Stomatognathic System - 21%, Germ Cells - 15%	Unclassified/Mixed - 24%, Stomatognathic System - 20%	Embryonic Structures - 21%	Skin - 41%, Musculoskeletal System - 20%, Female Genitalia - 17%, Hemic and Immune System - 17%	Exocrine Glands - 22%, Connective Tissue - 15%, Musculoskeletal System - 14%, Hemic and Immune Syste	Skin - 14%, Cardiovascular System - 11%	Male Genitalia - 15%, Endocrine System - 10%	Unclassified/Mixed - 19%, Pancreas - 14%, Female Genitalia - 10%	Liver - 16%, Endocrine System - 12%	Stomatognathic System - 54%, Musculoskeletal System - 18%	Hemic and Immune System - 100%	Nervous System - 100%	Pancreas - 100%	Unclassified/Mixed - 10%	Female Genitalia - 50%, Hemic and Immune System - 50%	Urinary Tract - 32%, Exocrine Glands - 14%, Cardiovascular System - 14%	Exocrine Glands - 31%, Urinary Tract - 31%, Digestive System - 23%	Sense Organs - 19%	Nervous System - 50%, Digestive System - 50%	Embryonic Structures - 29%, Nervous System - 24%, Connective Tissue - 19%	Female Genitalia - 16%, Respiratory System - 15%, Connective Tissue - 14%	Urinary Tract - 100%	Urinary Tract - 100%	Endocrine System - 32%, Skin - 25%, Connective Tissue - 12%, Cardiovascular System - 12%, Exocrine Gla	Digestive System - 24%, Skin - 21%, Liver - 17%	Digestive System - 100%	Urinary Tract - 25%, Pancreas - 22%, Male Genitalia - 14%	Nervous System - 100%	Skin · 24%, Embryonic Structures · 17%, Hemic and Immune System · 16%, Unclassified/Mixed · 16%	Musculoskeletal System - 53%, Sense Organs - 42%	Male Genitalia - 100%	Female Genitalia - 25%, Digestive System - 25%, Hemic and Immune System - 25%	Embryonic Structures - 12%, Endocrine System - 11%, Unclassified/Mixed - 10%	Stomatognathic System - 73%	Embryonic Structures - 100%	Embryonic Structures - 22%, Unclassified/Mixed - 20%, Urinary Tract - 19%	Embryonic Structures - 11%, Germ Cells - 11%
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SEQ	0 P	- 0	က	4	S.	9	7	æ	თ :	9	= :	12	4	9	8	ಜ		83	24	25	5 6	58	62	3	35	33	8	98	37	8	33	9	4	42	45	46	47	48	20	51

Template ID Tissue Distribution	dec Musculoskeletal System - 100%. 'dec Musculoskeletal System - 70%, Female Genitalia - 30%. 'dec Embryonic Structures - 34%, Connective Tissue - 24%, Liver - 18%. 'dec Urinary Tract - 33%, Nervous System - 25%, Digestive System - 25%. 'dec Skin - 59%, Endocrine System - 21%, Digestive System - 10%, Hemic and Immune System - 10%. 'dec Hemic and Immune System - 67%, Nervous System - 33%.	Cardiovascular System - 29%, Urinary Tract - 29%, Female Genitalia - 21% Skin - 25%, Endocrine System - 13%
Template ID	428745.2.dec 444839.17.dec 428362.36.dec 480710.12.dec 234137.10.dec 480630.4.dec 350399.5.dec 085713.2.dec	
SEQ ID NO	52 53 55 56 57 60 61	63

Table 4

Program	Description	Reference	Parameter Threshold
ABI FACTURA	A program that removes vector sequences and masks ambiguous bases in nucleic acid sequences.	PE Biosystems, Foster City, CA.	
ABIPARACEL FDF	A Fast Data Finder useful in comparing and annotating amino acid or nucleic acid sequences.	PE Biosystems, Foster City, CA; Paracel Inc., Pasadena, CA.	Mismatch <50%
ABI AutoAssembler	A program that assembles nucleic acid sequences.	PE Biosystems, Foster City, CA.	
BLAST	A Basic Local Alignment Search Tool useful in sequence similarity search for amino acid and nucleic acid sequences. BLAST includes five functions: blastp, blastn, tblastn, and tblastx.	Altschul, S.F. et al. (1990) J. Mol. Biol. 215:403-410; Altschul, S.F. et al. (1997) Nucleic Acids Res. 25:3389-3402.	ESTs: Probability value= 1.0E-8 or less Full Length sequences: Probability value= 1.0E-10 or less
FASTA	A Pearson and Lipman algorithm that searches for similarity between a query sequence and a group of sequences of the same type. FASTA comprises as least five functions: fasta, tfasta, fastx, tfastx, and ssearch.	Pearson, W.R. and D.J. Lipman (1988) Proc. Natl. Acad Sci. USA 85:2444-2448; Pearson, W.R. (1990) Methods Enzymol. 183:63-98; and Smith, T.F. and M.S. Waterman (1981) Adv. Appl. Math. 2:482-489.	ESTs: fasta E value=1.06E-6 Assembled ESTs: fasta Identity= 95% or greater and Match length=200 bases or greater; fastx E value=1.0E-8 or less Full Length sequences: fastx score=100 or greater
ВЫМРЅ	A BLocks IMProved Searcher that matches a sequence against those in BLOCKS, PRINTS, DOMO, PRODOM, and PFAM databases to search for gene families, sequence homology, and structural fingerprint regions.	Henikoff, S. and J.G. Henikoff (1991) Nucleic Acids Res. 19:6565-6572; Henikoff, J.G. and S. Henikoff (1996) Methods Enzymol. 266:88-105; and Attwood, T.K. et al. (1997) J. Chem. Inf. Comput. Sci. 37:417-424.	Score=1000 or greater; Ratio of Score/Strength = 0.75 or larger; and, if applicable, Probability value= 1.0E-3 or less
HMMER	An algorithm for searching a query sequence against hidden Markov model (HMM)-based databases of protein family consensus sequences, such as PFAM.	Krogh, A. et al. (1994) J. Mol. Biol. 235:1501-1531; Sonnhammer, E.L.L. et al. (1988) Nucleic Acids Res. 26:320-322.	Score=10-50 bits for PFAM hits, depending on individual protein families

Table 4 (cont.)

Parameter Threshold Normalized quality score > GCG- specified "HIGH" value for that particular Prosite motif. Generally, score=1.4-2.1.	Score= 120 or greater; Match length= 56 or greater	Score=3.5 or greater
Reference Gribskov, M. et al. (1988) CABIOS 4:61-66; Gribskov, M. et al. (1989) Methods Enzymol. 183:146-159; Bairoch, A. et al. (1997) Nucleic Acids Res. 25:217-221. Ewing, B. et al. (1998) Genome Res. 8:175-185; Ewing, B. and P. Green (1998) Genome Res. 8:175-185;	Smith, T.F. and M.S. Waterman (1981) Adv. Appl. Math. 2:482-489; Smith, T.F. and M.S. Waterman (1981) J. Mol. Biol. 147:195-197; and Green, P., University of Washington, Seattle, WA. Gordon, D. et al. (1998) Genome Res. 8:195-202.	Nielson, H. et al. (1997) Protein Engineering 10:1-6; Claverie, J.M. and S. Audic (1997) CABIOS 12:431-439. Bairoch, A. et al. (1997) Nucleic Acids Res. 25:217-221; Wisconsin Package Program Manual, version 9, page M51-59, Genetics Computer Group, Madison, WI.
An algorithm that searches for structural and sequence motifs in protein sequences that match sequence patterns defined in Prosite. A base-calling algorithm that examines automated sequencer traces with high sensitivity and probability.	A Phils Revised Assembly Program including SWAT and CrossMatch, programs based on efficient implementation of the Smith-Waterman algorithm, useful in searching sequence homology and assembling DNA sequences. A graphical tool for viewing and editing Phrap assemblies.	A weight matrix analysis program that scans protein sequences for the presence of secretory signal peptides. A program that searches amino acid sequences for patterns that matched those defined in Prosite.
Program ProfileScan Phred	Phrap Consed	SPScan Motifs

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CLAIMS

What is claimed is:

- An isolated polynucleotide comprising a polynucleotide sequence selected from the group
 consisting of:
 - a) a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63,
 - b) a naturally occurring polynucleotide sequence having at least 90% sequence identity to a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63,
 - c) a polynucleotide sequence complementary to a),
 - d) a polynucleotide sequence complementary to b), and
 - e) an RNA equivalent of a) through d).
 - 2. An isolated polynucleotide of claim 1, comprising a polynucleotide sequence selected from the group consisting of SEQ ID NO:1-63.
 - 3. An isolated polynucleotide comprising at least 60 contiguous nucleotides of a polynucleotide of claim 1.
- 4. A composition for the detection of expression of secretory polynucleotides comprising at least one of the polynucleotides of claim 1 and a detectable label.
 - 5. A method for detecting a target polynucleotide in a sample, said target polynucleotide having a sequence of a polynucleotide of claim 1, the method comprising:
- a) amplifying said target polynucleotide or fragment thereof using polymerase chain reaction amplification, and
 - b) detecting the presence or absence of said amplified target polynucleotide or fragment thereof, and, optionally, if present, the amount thereof.
- 6. A method for detecting a target polynucleotide in a sample, said target polynucleotide comprising a sequence of a polynucleotide of claim 1, the method comprising:
 - a) hybridizing the sample with a probe comprising at least 20 contiguous nucleotides comprising a sequence complementary to said target polynucleotide in the sample, and which probe specifically hybridizes to said target polynucleotide, under conditions whereby a hybridization complex is formed between said probe and said target polynucleotide or fragments thereof, and

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- b) detecting the presence or absence of said hybridization complex, and, optionally, if present, the amount thereof.
 - 7. A method of claim 5, wherein the probe comprises at least 30 contiguous nucleotides.
 - 8. A method of claim 5, wherein the probe comprises at least 60 contiguous nucleotides.
- 9. A recombinant polynucleotide comprising a promoter sequence operably linked to a polynucleotide of claim 1.
 - 10. A cell transformed with a recombinant polynucleotide of claim 9.
 - 11. A transgenic organism comprising a recombinant polynucleotide of claim 9.
- 12. A method for producing a secretory polypeptide, the method comprising:
 - a) culturing a cell under conditions suitable for expression of the secretory polypeptide, wherein said cell is transformed with a recombinant polynucleotide of claim 9, and
 - b) recovering the secretory polypeptide so expressed.
- 20 13. A purified secretory polypeptide encoded by at least one of the polynucleotides of claim 2.
 - 14. An isolated antibody which specifically binds to a secretory polypeptide of claim 13.
- 15. A method of identifying a test compound which specifically binds to the secretory polypeptide of claim 13, the method comprising the steps of:
 - a) providing a test compound;
 - b) combining the secretory polypeptide with the test compound for a sufficient time and under suitable conditions for binding; and
- c) detecting binding of the secretory polypeptide to the test compound, thereby identifying the test compound which specifically binds the secretory polypeptide.
 - 16. A microarray wherein at least one element of the microarray is a polynucleotide of claim 3.

17. A method for generating a transcript image of a sample which contains polynucleotides, the method comprising the steps of:

a) labeling the polynucleotides of the sample,

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- b) contacting the elements of the microarray of claim 16 with the labeled polynucleotides of the sample under conditions suitable for the formation of a hybridization complex, and
 - c) quantifying the expression of the polynucleotides in the sample.
- 18. A method for screening a compound for effectiveness in altering expression of a target polynucleotide, wherein said target polynucleotide comprises a polynucleotide sequence of claim 1, the method comprising:
- a) exposing a sample comprising the target polynucleotide to a compound, under conditions suitable for the expression of the target polynucleotide,
 - b) detecting altered expression of the target polynucleotide, and
- c) comparing the expression of the target polynucleotide in the presence of varying amounts of
 the compound and in the absence of the compound.
 - 19. A method for assessing toxicity of a test compound, said method comprising:
 - a) treating a biological sample containing nucleic acids with the test compound;
- b) hybridizing the nucleic acids of the treated biological sample with a probe comprising at
 least 20 contiguous nucleotides of a polynucleotide of claim 1 under conditions whereby a specific hybridization complex is formed between said probe and a target polynucleotide in the biological sample, said target polynucleotide comprising a polynucleotide sequence of a polynucleotide of claim 1 or fragment thereof;
 - c) quantifying the amount of hybridization complex; and
- d) comparing the amount of hybridization complex in the treated biological sample with the amount of hybridization complex in an untreated biological sample, wherein a difference in the amount of hybridization complex in the treated biological sample is indicative of toxicity of the test compound.

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SEQUENCE LISTING

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